

GenCore version 5.1.4_p5_4578
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OM nucleic - nucleic search, using sw model

Run on: May 25, 2003, 03:08:07 ; Search time 234 Seconds
(without alignments)
1761.178 Million cell updates/sec

Title: US-09-660-568-49

Perfect score: 183
Sequence: 1 cacacactccccattctga.....ctgtctctcggagctcaacca 183

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 1125999159 residues

total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

- 1: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1980.DAT.*
- 2: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1981.DAT.*
- 3: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1982.DAT.*
- 4: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1983.DAT.*
- 5: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1984.DAT.*
- 6: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1985.DAT.*
- 7: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1986.DAT.*
- 8: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1987.DAT.*
- 9: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1988.DAT.*
- 10: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1989.DAT.*
- 11: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1990.DAT.*
- 12: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1991.DAT.*
- 13: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1992.DAT.*
- 14: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1993.DAT.*
- 15: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1994.DAT.*
- 16: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1995.DAT.*
- 17: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1996.DAT.*
- 18: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1997.DAT.*
- 19: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1998.DAT.*
- 20: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1999.DAT.*
- 21: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2000.DAT.*
- 22: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT.*
- 23: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT.*
- 24: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2002.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	183	100.0	183	19	AAV38517
2	183	100.0	183	20	AAV38517
3	41	22.4	2113	23	AAV38517
4	35.2	19.2	164	22	AAV38517
5	35.2	19.2	164	22	AAV38517
6	35.2	19.2	164	22	AAV38517
7	35.2	19.2	164	22	AAV38517
8	35.2	19.2	164	22	AAV38517
9	35.2	19.2	164	22	AAV38517

10	33.6	18.4	1524	23	ABL30203	Drosophila melanog
11	32.8	17.9	3819	23	ABL30202	Drosophila melanog
12	31.2	17.0	3698	22	AAK73760	Human immune/haema
13	31.2	17.0	3698	22	AAK83893	Human immune/haema
14	31.2	17.0	10548	22	AAK64703	Human immune/haema
15	31.2	17.0	10548	22	AAK69813	Human immune/haema
16	31.2	17.0	10548	22	AAK73759	Human immune/haema
17	31.2	17.0	10548	22	AAK80461	Human immune/haema
18	31.2	17.0	10548	22	AAK83894	Human immune/haema
19	30.8	16.8	5616	22	AAK79614	Human immune/haema
20	30.6	16.7	3387	23	AAK91234	DNA encoding novel
21	30.6	16.7	5216	22	AAK28261	Genomic sequence #
22	30.6	16.7	5216	22	AAK31528	Human DNA for a no
23	30.6	16.7	5216	24	ABQ66852	Human polynucleoti
24	30.2	16.5	433	22	AAI89606	Human polynucleoti
25	29.8	16.3	13852	22	ABA17744	Human nervous syst
26	29.8	16.3	13852	22	ABA17745	Human nervous syst
27	29.8	16.3	13852	22	ABA21395	Human nervous syst
28	29.8	16.3	13852	22	ABA21396	Human nervous syst
29	29.6	16.2	4068	22	AAI86707	Human polynucleoti
30	29.2	16.0	2086	23	ABL12837	Drosophila melanog
31	29.2	16.0	3598	22	AAI64774	Human glucose tran
32	29.2	16.0	18733	22	AAK80682	Human immune/haema
33	29.2	16.0	100301	24	AEQ88176	Human osteoblast d
34	29	15.8	3212	24	ABK11142	CDNA of human clon
35	29	15.8	38886	20	AAI23897	Murine LOBO homolo
36	29	15.8	49999	20	AAI23891	Murine LOBO homolo
37	28.8	15.7	431	21	AAC28387	Human secreted pro
38	28.8	15.7	1090	18	AAI72790	Metastasis inducin
39	28.8	15.7	894	23	ABK42719	Genomic sequence #
40	28.4	15.5	1089	24	ABK63947	CDNA encoding huma
41	28.4	15.5	2001	24	AAI6204	Human disulfide co
42	28.4	15.5	2513	24	AAI6206	Human disulfide co
43	28.4	15.5	44576	21	AAI61522	Cosmid CVO14 conta
44	28.2	15.4	341	22	ABA16249	Human nervous syst
45	28.2	15.4	1907	20	AAV84508	Human secreted pro

ALIGNMENTS

RESULT 1

AAV38517 standard; DNA; 183 BP.

AAV38517;

08-OCT-1998 (first entry)

DNA marker of metastatic prostate cancer, UC Band#321.

DNA marker; metastatic prostate cancer; human; UC Band#321; detection;

disease marker identification; lupus erythematosus; rheumatoid arthritis;

multiple sclerosis; asthma; myasthenia gravis; autoimmune thyroiditis;

amyloid lateral sclerosis; interstitial cystitis; prostatitis; ss.

Homo sapiens.

WO9824935-A1.

11-JUN-1998.

05-DEC-1997; 97WO-US22105.

24-MAR-1997; 97US-0041576.

06-DEC-1996; 96US-0032619.

12-DEC-1996; 96US-0032701.

(UROC-) UROCOR INC.

An G, O'Hara M, Ralph D, Veltri R;

WPI; 1998-333350/29.

XX Identifying markers for disease states - by amplifying RNA from
 PT peripheral blood and identifying RNA which is differential expressed
 PT between normal and disease state subjects
 XX
 PS Claim 17; Page 92; 158pp; English.
 XX
 CC This sequence represents a DNA marker of metastatic prostate cancer,
 CC designated UC Band#321, and was identified using a method of the
 CC invention. The method is for identifying markers for a disease state, and
 CC comprises: (a) providing a first set of peripheral blood mRNAs from one
 CC or more subjects known to exhibit the disease state and a second set of
 CC peripheral blood mRNAs from one or more normal subjects; (b) amplifying
 CC both sets of mRNAs to provide nucleic acid amplification products;
 CC (c) comparing the sets of amplification products; and (d) identifying
 CC those mRNAs that are differentially expressed between normal subjects and
 CC subjects exhibiting the disease state; where a difference in quantity of
 CC expression of an mRNA is indicative of a disease marker. This marker
 CC sequence can be used in a method of detecting a metastatic cancer disease
 CC state, especially for detection prostate cancer. Using the methods, a
 CC disease state may be detected, diagnosed, or a prognosis may be delivered
 CC by examining a blood sample rather than relying on a more invasive, or
 CC less sensitive test. In addition, a subject may be monitored for disease
 CC progression, status and response to therapies through monitoring of
 CC differentially expressed disease markers. The methods can be used for
 CC diseases such as cancer (especially metastatic or prostate cancer),
 CC asthma, lupus erythematosus, rheumatoid arthritis, multiple sclerosis,
 CC myasthenia gravis, autoimmune thyroiditis, amyloid lateral sclerosis,
 CC interstitial cystitis, prostatitis or other systemic or chronic conditions.
 XX
 SQ Sequence 183 BP; 43 A; 52 C; 45 G; 43 T; 0 other;
 Query Match 100.0%; Score 183; DB 19; Length 183;
 Best Local Similarity 100.0%; Pred. No. 2.8e-53;
 Matches 183; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CACACACTCCCCATCTGAGGCCCAAGAGGCTCATCCCTAAGGATGCCAGAGATCCAA 60
 DB 1 CACACACTCCCCATCTGAGGCCCAAGAGGCTCATCCCTAAGGATGCCAGAGATCCAA 60
 QY 61 GTGCAGAGGAGAAATGTGGTGGAGGCTATTATTATCCCCAGTGCCTTCCCTGGGCTAT 120
 DB 61 GTGCAGAGGAGAAATGTGGTGGAGGCTATTATTATCCCCAGTGCCTTCCCTGGGCTAT 120
 QY 121 GGATGAACAGTGCGTGACTTCATCTAGGAAAGAGCTATGGCTTCTGCTCTCTGGAGCTCA 180
 DB 121 GGATGAACAGTGCGTGACTTCATCTAGGAAAGAGCTATGGCTTCTGCTCTCTGGAGCTCA 180
 181 CCA 183
 181 CCA 183
 RESULT 2
 AAZ30719
 ID AAZ30719 standard; cDNA; 183 BP.
 XX
 AC AAZ30719;
 XX
 DT 19-JAN-2000 (first entry)
 XX
 XX Human UC Band #321 cDNA.
 XX
 KW UC Band #321; marker; expression; diagnosis;
 KW differential; disease; cancer; metastatic; breast cancer; prostate;
 KW peripheral leukocyte; immune response; asthma; lupus erythematosus;
 KW rheumatoid arthritis; multiple sclerosis; myasthenia gravis;
 KW autoimmune thyroiditis; amyotrophic lateral sclerosis; ALS;
 KW interstitial cystitis; prostatitis; mRNA; reverse transcriptase PCR;
 KW RT-PCR; screening; early; diagnosis; prognosis; monitoring; ss.
 OS
 XX Homo sapiens.
 XX

PN WO9949083-A1.
 XX
 PD 30-SEP-1999.
 XX
 PF 24-MAR-1999; 99WO-US06488.
 XX
 PR 24-MAR-1998; 98US-0046894.
 XX
 PA (UROC-) UROCOR INC.
 XX
 PI Ralph D, An G, O'Hara SM, Veltri RW;
 XX
 DR WPI; 1999-591105/50.
 XX
 PT Identifying markers of human disease, specifically for diagnosis of
 PT metastatic prostatic and breast cancers
 XX
 PS Claim 17; Page 124-125; 225pp; English.
 XX
 CC This sequence represents a human cDNA sequence designated UC Band #321,
 CC which encodes a previously undescribed gene product. The expression of
 CC this gene in peripheral leukocytes was examined using reverse
 CC transcriptase-PCR (RT-PCR) primers AAZ30739-230740. This gene was found
 CC to be differentially expressed between healthy subjects and patients
 CC with metastatic cancers (especially those of the prostate or breast) and
 CC may therefore be used as a marker for such diseases. Detecting
 CC levels of such human disease markers is used for diagnosis (also
 CC prognosis and monitoring) of diseases, including metastatic or
 CC organ-confined cancers, and diseases which also elicit an immune
 CC response such as asthma, lupus erythematosus, rheumatoid arthritis,
 CC multiple sclerosis, myasthenia gravis, autoimmune thyroiditis,
 CC amyotrophic lateral sclerosis (ALS), interstitial cystitis and
 CC prostatitis, but especially metastatic prostatic and breast cancer. A
 CC particular use is differentiating between prostatic cancer and benign
 CC prostatic hypertrophy, and between advanced and localised prostatic
 CC cancer. By multivariate analysis of several different markers. Cancers
 CC can be treated by administering sequences antisense to sequences that
 CC encode human disease markers. This method detects a leukocyte response
 CC to disease rather than products of diseased cells, so is suitable for
 CC large-scale screening of asymptomatic subjects. Disease can be detected
 CC at an early stage, when few, if any, diseased cells are present in the
 CC circulation. Analysis of blood samples eliminates the need for more
 CC invasive methods for obtaining samples.
 XX
 SQ Sequence 183 BP; 43 A; 52 C; 45 G; 43 T; 0 other;
 Query Match 100.0%; Score 183; DB 20; Length 183;
 Best Local Similarity 100.0%; Pred. No. 2.8e-53;
 Matches 183; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CACACACTCCCCATCTGAGGCCCAAGAGGCTCATCCCTAAGGATGCCAGAGATCCAA 60
 DB 1 CACACACTCCCCATCTGAGGCCCAAGAGGCTCATCCCTAAGGATGCCAGAGATCCAA 60
 QY 61 GTGCAGAGGAGAAATGTGGTGGAGGCTATTATTATCCCCAGTGCCTTCCCTGGGCTAT 120
 DB 61 GTGCAGAGGAGAAATGTGGTGGAGGCTATTATTATCCCCAGTGCCTTCCCTGGGCTAT 120
 QY 121 GGATGAACAGTGCGTGACTTCATCTAGGAAAGAGCTATGGCTTCTGCTCTCTGGAGCTCA 180
 DB 121 GGATGAACAGTGCGTGACTTCATCTAGGAAAGAGCTATGGCTTCTGCTCTCTGGAGCTCA 180
 181 CCA 183
 181 CCA 183
 RESULT 3
 AAS77249
 ID AAS77249 standard; cDNA; 2113 BP.
 XX
 AC AAS77249;
 XX

DT 13-FEB-2002 (first entry)
 XX DNA encoding novel human diagnostic protein #13053.
 XX Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
 KW Homo sapiens.
 XX WO200175067-A2.
 XX 11-OCT-2001.
 XX 30-MAR-2001; 2001WO-US08631.
 XX 31-MAR-2000; 2000US-0540217.
 PR 23-AUG-2000; 2000US-0649167.
 XX (HYSE-) HYSEQ INC.
 PA Drmanac RT, Liu C, Tang YT;
 XX WPI; 2001-639362/73.
 DR P-PSDB; ABG13062.
 XX New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 responsible for genetic disorders or other traits and to assess
 PT biodiversity.
 XX Claim 1; SEQ ID No 13053; 103pp; English.
 XX The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.
 XX Sequence 2113 BP; 761 A; 474 C; 421 G; 457 T; 0 other;
 Query Match 22.4%; Score 41; DB 23; Length 2113;
 Best Local Similarity 64.2%; Pred. No. 0.00086;
 Matches 79; Conservative 0; Mismatches 40; Indels 4; Gaps 1;

QY 30 GGCTCATCCCTTAGGATGTCAGAGATCCAGTCCAGAGAGAGATGTTGGAGCTATT 89
 III III I II III III III III III III III III III
 Db 170 GCCCAATCGACGACCTCGAAGGAGATCAAAAGGGTAGAAGGAGA---GATTTCAGGTATC 225
 QY 90 TATTCCCCCAGTGCCTCCCTGCTGGGCTATGGATGAACAGTGGCTGACTTCATCTAGGA 149
 T III III III III III III III III III III III III
 Db 226 TGTTCCTCCCTGGTCACTTCCCTGCTGGGCTATGGCTTGGCACTGGCTGGGATCCCTTACCT 285
 QY 150 AAG 152
 III
 Db 286 AAG 288

RESULT 4
 ABA51400
 ID ABA51400 standard; DNA; 164 BP.
 XX AC ABA51400;
 XX 01-FEB-2002 (first entry)
 DT Human breast cell single exon nucleic acid probe #10095.
 XX Human; microarray; single exon probe; gene expression; breast;
 KW disease; cancer; ss.
 XX Homo sapiens.
 OS WO200157271-A2.
 XX 09-AUG-2001.
 PD 30-JAN-2001; 2001WO-US00662.
 PF 04-FEB-2000; 2000US-0180312.
 XX 26-MAY-2000; 2000US-0207456.
 PR 30-JUN-2000; 2000US-0608408.
 PR 03-AUG-2000; 2000US-0632366.
 PR 21-SEP-2000; 2000US-0234687.
 PR 27-SEP-2000; 2000US-0236359.
 PR 04-OCT-2000; 2000GB-0024263.
 XX (MOLE-) MOLECULAR DYNAMICS INC.
 PA Penn SG, Hanzel DK, Chen W, Rank DR;
 XX WPI; 2001-496933/54.
 DR New spatially-addressable set of single exon nucleic acid probes,
 PT useful for measuring gene expression in sample derived from human
 PT breast, comprises number of single exon nucleic acid probes -
 PS Claim 4; SEQ ID NO 10095; 327pp + sequence listing; English.
 XX The invention relates to a spatially-addressable set of single exon
 CC nucleic acid probes for measuring gene expression in a sample derived
 CC from human breast and BT 474 cells. The method involves contacting
 CC the probes with a collection of detectably labelled nucleic acids
 CC derived from mRNA of human breast, and then measuring the label
 CC bound to each probe of the microarray. The probes are useful for
 CC verifying the expression of regions of genomic DNA predicted to
 CC encode proteins. They are useful for gene discovery, and for
 CC determining predisposition and/or prognosing breast disease. Gene
 CC expression analysis is useful for assessing the toxicity of chemical
 CC agents on cells. The microarray of this invention presents a far greater
 CC diversity of probes for measuring gene expression, with far less bias
 CC than expressed sequence tag microarrays. The method is suitable for
 CC rapid production of functional information from genomic sequence. The
 CC present sequence is a single exon nucleic acid probe of the invention.
 CC Note: The sequence data for this patent did not form part of the
 CC printed specification, but was obtained in electronic format directly
 CC from WIPO at ftp.wipo.int/pub/published_pct_sequences.
 XX Sequence 164 BP; 39 A; 39 C; 44 G; 42 T; 0 other;
 Query Match 19.2%; Score 35.2; DB 22; Length 164;
 Best Local Similarity 71.9%; Pred. No. 0.031;
 Matches 46; Conservative 0; Mismatches 18; Indels 0; Gaps 0;

QY 52 GAGATCCAGTGCAGAGAGAGATGTGTGAGGCTATTATTTCCCGGCTTCCCTG 111
 III III III III III III III III III III III III
 Db 100 GAGATCAGAGACGACAGAGAGAGATGAAGTGAGGATATTTATTTCCCTTGGCTCTCTCTTG 159
 QY 112 CTGG 115
 II
 Db 160 TGGG 163

RESULT 5

ABA69411
ID ABA69411 standard; DNA; 164 BP.
XX AC
XX ABA69411;
XX
DT 01-FEB-2002 (first entry)
XX
DE Human foetal liver single exon nucleic acid probe #17716.
XX
KW Human; foetal liver; gene expression; single exon nucleic acid probe; ss.
XX
XX Homo sapiens.
OS
XX
XX WO200157277-A2.
PN
XX
PD 09-AUG-2001.
XX
PF 30-JAN-2001; 2001WO-US00669.
XX
PR 04-FEB-2000; 2000US-0180312.
PR 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
PR 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
PR 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024263.
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
XX Penn SG, Hanzel DK, Chen W, Rank DR;
PI WPI; 2001-483447/52.
XX
XX Human genome-derived single exon nucleic acid probes useful for
PT analyzing gene expression in human fetal liver -
XX
XX Claim 4; SEQ ID NO 17716; 639pp + sequence listing; English.
PS
XX The invention relates to a single exon nucleic acid probe for
CC measuring human gene expression in a sample derived from human foetal
CC liver. The single exon nucleic acid probes may be used for predicting,
CC measuring and displaying gene expression in samples derived from human
CC fetal liver. The present sequence is a single exon nucleic acid
CC probe of the invention.
CC Note: The sequence data for this patent did not form part of the
CC printed specification, but was obtained in electronic format directly
CC from WIPO at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 164 BP; 39 A; 39 C; 44 G; 42 T; 0 other;

Query Match 19.2%; Score 35.2; DB 22; Length 164;
Best Local Similarity 71.9%; Pred. No. 0.031;
Matches 46; Conservative 0; Mismatches 18; Indels 0; Gaps 0;

QY 52 GAGATCAAGTCGAGAGAGAGATGTGGTGGAGCTATTATTCCTCCAGTGCTTCCCTG 111
||||| ||||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||
DB 100 GAGATCAGAGCAGCAGAGAGAGATGAAGTCAGGATATTTATTCCTTGGCTCTCTCTTG 159
QY 112 CTGG 115
DB 160 TGGG 163

RESULT 6

ABA36346
ID ABA36346 standard; DNA; 164 BP.
XX AC
XX ABA36346;
XX
DT 23-JAN-2002 (first entry)

XX

DE Probe #14812 for gene expression analysis in human heart cell sample.
XX
XX Human; gene expression; heart; microarray; vascular system; probe;
KW cardiovascular disease; hypertension; cardiac arrhythmia;
KW congenital heart disease; ss.
XX
XX Homo sapiens.
OS
XX
XX WO200157274-A2.
PN
XX
PD 09-AUG-2001.
XX
PF 30-JAN-2001; 2001WO-US00666.
XX
PR 04-FEB-2000; 2000US-0180312.
PR 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
PR 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
PR 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024263.
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
XX Penn SG, Hanzel DK, Chen W, Rank DR;
PI WPI; 2001-488899/53.
XX
XX Single exon nucleic acid probes for analyzing gene expression in human
PT hearts -
PT
XX

PS Claim 4; SEQ ID No 14812; 530pp; English.
XX
XX The present invention relates to single exon nucleic acid probes for
CC measuring human gene expression in a sample derived from human heart. The
CC present sequence is one such probe. The probes may be used for
CC predicting, measuring and displaying gene expression in samples derived
CC from the human heart via microarrays. By measuring gene expression, the
CC probes are useful for predicting, diagnosing, grading, staging,
CC monitoring and prognosing diseases of the human heart and vascular system
CC e.g. cardiovascular disease, hypertension, cardiac arrhythmias and
CC congenital heart disease.
CC Note: The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 164 BP; 39 A; 39 C; 44 G; 42 T; 0 other;

Query Match 19.2%; Score 35.2; DB 22; Length 164;
Best Local Similarity 71.9%; Pred. No. 0.031;
Matches 46; Conservative 0; Mismatches 18; Indels 0; Gaps 0;

QY 52 GAGATCAAGTCGAGAGAGAGATGTGGTGGAGCTATTATTCCTCCAGTGCTTCCCTG 111
||||| ||||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||
DB 100 GAGATCAGAGCAGCAGAGAGAGATGAAGTCAGGATATTTATTCCTTGGCTCTCTCTTG 159
QY 112 CTGG 115
DB 160 TGGG 163

RESULT 7

AAK17679
ID AAK17679 standard; DNA; 164 BP.
XX
XX AAK17679;
XX
XX
DT 05-NOV-2001 (first entry)
XX
DE Human brain expressed single exon probe SEQ ID NO: 17670.
XX
KW Human; brain expressed exon; gene expression analysis; probe;

```
KW microarray; Alzheimer's disease; multiple sclerosis; schizophrenia;
KW epilepsy; cancer; ss.
XX Homo sapiens.
XX WO200157275-A2.
XX 09-AUG-2001.
XX 30-JAN-2001; 2001WO-US00667.
XX 04-FEB-2000; 2000US-0180312.
XX 26-MAY-2000; 2000US-0207456.
XX 30-JUN-2000; 2000US-0608408.
XX 03-AUG-2000; 2000US-0632366.
XX 21-SEP-2000; 2000US-0234687.
XX 27-SEP-2000; 2000US-0236359.
XX 04-OCT-2000; 2000GB-0024263.
XX (MOLE-) MOLECULAR DYNAMICS INC.
XX PA Penn SG, Hanzel DK, Chen W, Rank DR;
XX WPI; 2001-483446/52.
XX Single exon nucleic acid probes for analyzing gene expression in human
XX brains -
XX Example 4; SEQ ID NO: 17670; 650pp + Sequence Listing; English.
XX The present invention provides a number of single exon nucleic acid
XX probes which are derived from genomic sequences expressed in the human
XX brain. They can be used to measure gene expression in brain cell samples,
XX which may enable the diagnosis and improved treatment of nervous system
XX diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia,
XX epilepsy and cancers. The present sequence is one of the probes of the
XX invention.
XX SQ Sequence 164 BP; 39 A; 39 C; 44 G; 42 T; 0 other;
XX Query Match 19.2%; Score 35.2; DB 22; Length 164;
XX Best Local Similarity 71.9%; Pred. No. 0.031;
XX Matches 46; Conservative 0; Mismatches 18; Indels 0; Gaps 0;
Oy 52 GAGATCCAGTGCGAGAGGAGAAATGTGTGAGGCTATTATTTCCCCAGTCCCTTCCCTG 111
Db 100 GAGATCAGAGAGCAGAGAGAGAAATGTGTGAGGCTATTATTTCCCTTGGCTCTCTCTTG 159
Oy 112 CTGG 115
Db 160 TGGG 163
XX RESULT 8
XX AAK43494
XX ID AAK43494 standard; DNA; 164 BP.
XX AC AAK43494;
XX 06-NOV-2001 (first entry)
XX Human bone marrow expressed single exon probe SEQ ID NO: 18051.
XX Human; bone marrow expressed exon; gene expression analysis; probe;
XX microarray; cancer; leukaemia; lymphoma; myeloma; ss.
XX Homo sapiens.
XX WO200157276-A2.
XX 09-AUG-2001.
XX 30-JAN-2001; 2001WO-US00668.
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XX 04-FEB-2000; 2000US-0180312.
XX 26-MAY-2000; 2000US-0207456.
XX 30-JUN-2000; 2000US-0608408.
XX 03-AUG-2000; 2000US-0632366.
XX 21-SEP-2000; 2000US-0234687.
XX 27-SEP-2000; 2000US-0236359.
XX 04-OCT-2000; 2000GB-0024263.
XX (MOLE-) MOLECULAR DYNAMICS INC.
XX PA Penn SG, Hanzel DK, Chen W, Rank DR;
XX WPI; 2001-488900/53.
XX Human genome-derived single exon nucleic acid probes useful for
XX analyzing gene expression in human bone marrow -
XX Example 4; SEQ ID NO: 18051; 658pp + Sequence Listing; English.
XX The present invention provides a number of single exon nucleic acid
XX probes which are derived from genomic sequences expressed in the human
XX bone marrow. They can be used to measure gene expression in bone marrow
XX samples, which may enable the improved diagnosis and treatment of cancers
XX such as lymphoma, leukaemia and myeloma. The present sequence is one of
XX the probes of the invention.
XX SQ Sequence 164 BP; 39 A; 39 C; 44 G; 42 T; 0 other;
XX Query Match 19.2%; Score 35.2; DB 22; Length 164;
XX Best Local Similarity 71.9%; Pred. No. 0.031;
XX Matches 46; Conservative 0; Mismatches 18; Indels 0; Gaps 0;
Oy 52 GAGATCCAGTGCGAGAGGAGAAATGTGTGAGGCTATTATTTCCCCAGTCCCTTCCCTG 111
Db 100 GAGATCAGAGAGCAGAGAGAGAAATGTGTGAGGCTATTATTTCCCTTGGCTCTCTCTTG 159
Oy 112 CTGG 115
Db 160 TGGG 163
XX RESULT 9
XX AAI24280
XX ID AAI24280 standard; DNA; 164 BP.
XX AC AAI24280;
XX 12-OCT-2001 (first entry)
XX Probe #14213 for gene expression analysis in human cervical cell sample.
XX Probe; human; microarray; gene expression; cervical epithelial cell;
XX cervical cancer; ss.
XX Homo sapiens.
XX WO200157278-A2.
XX 09-AUG-2001.
XX 30-JAN-2001; 2001WO-US00670.
XX 04-FEB-2000; 2000US-0180312.
XX 26-MAY-2000; 2000US-0207456.
XX 30-JUN-2000; 2000US-0608408.
XX 03-AUG-2000; 2000US-0632366.
XX 21-SEP-2000; 2000US-0234687.
XX 27-SEP-2000; 2000US-0236359.
XX 04-OCT-2000; 2000GB-0024263.
XX (MOLE-) MOLECULAR DYNAMICS INC.
XX PA
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PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX WPI; 2001-488901/53.
XX Human genome-derived single exon nucleic acid probes useful for
PT analyzing gene expression in human cervical epithelial cells -
PS Claim 25; SEQ ID No 14213; 487pp; English.
XX The present invention relates to human single exon nucleic acid probes
CC (SENP). The present sequence is one such probe. The SENPs are derived
CC from human Hela cells. The SENPs can be used to produce a single exon
CC microarray, which can be used for measuring human gene expression in a
CC sample derived from human cervical epithelial cells. By measuring gene
CC expression, the probes are therefore useful in grading and/or staging
CC of diseases of the cervix, notably cervical cancer.
CC Note: The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX Sequence 164 BP; 39 A; 39 C; 44 G; 42 T; 0 other;
Query Match 19.2%; Score 35.2; DB 22; Length 164;
Best Local Similarity 71.9%; Pred. No. 0.031;
Matches 46; Conservative 0; Mismatches 18; Indels 0; Gaps 0;
QY 52 GAGATCCAAAGTCGACGAGAGAGATGGTGAGGCTATTATTCCCGCTTCCCTG 111
DB 100 GAGATCAGAGCAGCAGAGAGAGATGAAGTGAGGATATTATTCCCTTGGCTCTCTCTG 159
QY 112 CTGG 115
DB 160 TGGG 163
RESULT 10
ABL30203
ID ABL30203 standard; DNA; 1524 BP.
XX ABL30203;
AC ABL30203;
DT 26-MAR-2002 (first entry)
DE Drosophila melanogaster genomic polynucleotide SEQ ID NO 42082.
XX Drosophila; developmental biology; cell signalling; insecticide;
KW pharmaceutical; gene; ds.
XX Drosophila melanogaster.
PN WO200171042-A2.
XX 27-SEP-2001.
XX 23-MAR-2001; 2001WO-US09231.
XX 23-MAR-2000; 2000US-191637P.
PR 11-JUL-2000; 2000US-0614150.
XX (PEKE) PE CORP NY.
XX Venter JC, Adams M, Li PWD, Myers EW;
XX WPI; 2001-656860/75.
XX New isolated nucleic acid detection reagent for detecting 1000 or more
PT genes from Drosophila and for elucidating cell signalling and cell-cell
PT interactions -
XX Claim 1; SEQ ID NO 42082; 21pp + Sequence Listing; English.
XX The invention relates to an isolated nucleic acid detection reagent
CC capable of detecting 1000 or more genes from Drosophila. The invention is

CC useful in developmental biology and in elucidating cell signalling and
CC cell-cell interactions in higher eukaryotes for the development of
CC insecticides, therapeutics and pharmaceutical drugs. The invention
CC discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA
CC sequences (ABL01840-ABL16175) and the encoded proteins
CC (ABB57737-ABB72072).
CC The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX Sequence 1524 BP; 455 A; 333 C; 353 G; 383 T; 0 other;
Query Match 18.4%; Score 33.6; DB 23; Length 1524;
Best Local Similarity 53.9%; Pred. No. 0.27;
Matches 69; Conservative 0; Mismatches 59; Indels 0; Gaps 0;
QY 38 CTAAGGATGTCACAGATCCCAAGTCGACGAGAGAGATGGTGAGGCTATTATTCC 97
DB 400 CCAACAGATTGTCAGATGTCGGGCGAGGTCACAGAAAGTGGTGATCATTTAATTCACC 459
QY 98 CAGTGCCTTCCTGCTGGGCTATGATGAACAGTGGCTGACTTCATCTAGGAAAGAGCTA 157
DB 460 ATTGCTGCACAGCAGGCAAGGCCAAGGATGTCCACAGGAAGTGATATCCAGCAAAACTCA 519
QY 158 TGGCTTCT 165
DB 520 GAGTTTAT 527
RESULT 11
ABL30202/c
ID ABL30202 standard; DNA; 3819 BP.
XX ABL30202;
AC ABL30202;
DT 26-MAR-2002 (first entry)
DE Drosophila melanogaster genomic polynucleotide SEQ ID NO 42079.
XX Drosophila; developmental biology; cell signalling; insecticide;
KW pharmaceutical; gene; ds.
XX Drosophila melanogaster.
PN WO200171042-A2.
XX 27-SEP-2001.
XX 23-MAR-2001; 2001WO-US09231.
PR 23-MAR-2000; 2000US-191637P.
PR 11-JUL-2000; 2000US-0614150.
XX (PEKE) PE CORP NY.
XX Venter JC, Adams M, Li PWD, Myers EW;
XX WPI; 2001-656860/75.
XX New isolated nucleic acid detection reagent for detecting 1000 or more
PT genes from Drosophila and for elucidating cell signalling and cell-cell
PT interactions -
XX Claim 1; SEQ ID NO 42079; 21pp + Sequence Listing; English.
XX The invention relates to an isolated nucleic acid detection reagent
CC capable of detecting 1000 or more genes from Drosophila. The invention is
CC useful in developmental biology and in elucidating cell signalling and
CC cell-cell interactions in higher eukaryotes for the development of
CC insecticides, therapeutics and pharmaceutical drugs. The invention
CC discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA
CC sequences (ABL01840-ABL16175) and the encoded proteins
CC (ABB57737-ABB72072).

PR	08-SEP-2000;	2000US-0232080;
PR	08-SEP-2000;	2000US-0232081;
PR	12-SEP-2000;	2000US-0232168;
PR	14-SEP-2000;	2000US-0232169;
PR	14-SEP-2000;	2000US-0232397;
PR	14-SEP-2000;	2000US-0232398;
PR	14-SEP-2000;	2000US-0232399;
PR	14-SEP-2000;	2000US-0232400;
PR	14-SEP-2000;	2000US-0232401;
PR	14-SEP-2000;	2000US-0233063;
PR	14-SEP-2000;	2000US-0233064;
PR	14-SEP-2000;	2000US-0233065;
PR	21-SEP-2000;	2000US-0233233;
PR	21-SEP-2000;	2000US-0234223;
PR	21-SEP-2000;	2000US-0234274;
PR	25-SEP-2000;	2000US-0234997;
PR	25-SEP-2000;	2000US-0234998;
PR	26-SEP-2000;	2000US-0234984;
PR	27-SEP-2000;	2000US-0235834;
PR	27-SEP-2000;	2000US-0235835;
PR	29-SEP-2000;	2000US-0236327;
PR	29-SEP-2000;	2000US-0236327;
PR	29-SEP-2000;	2000US-0236368;
PR	29-SEP-2000;	2000US-0236369;
PR	29-SEP-2000;	2000US-0236370;
PR	02-OCT-2000;	2000US-0236802;
PR	02-OCT-2000;	2000US-0237037;
PR	02-OCT-2000;	2000US-0237038;
PR	02-OCT-2000;	2000US-0237039;
PR	02-OCT-2000;	2000US-0237040;
PR	13-OCT-2000;	2000US-0239935;
PR	13-OCT-2000;	2000US-0239937;
PR	20-OCT-2000;	2000US-0241261;
PR	20-OCT-2000;	2000US-0241262;
PR	20-OCT-2000;	2000US-0241281;
PR	20-OCT-2000;	2000US-0241785;
PR	20-OCT-2000;	2000US-0241786;
PR	20-OCT-2000;	2000US-0241787;
PR	20-OCT-2000;	2000US-0241808;
PR	20-OCT-2000;	2000US-0241809;
PR	20-OCT-2000;	2000US-0241826;
PR	01-NOV-2000;	2000US-0244617;
PR	08-NOV-2000;	2000US-0245674;
PR	08-NOV-2000;	2000US-0245675;
PR	08-NOV-2000;	2000US-0246476;
PR	08-NOV-2000;	2000US-0246477;
PR	08-NOV-2000;	2000US-0246478;
PR	08-NOV-2000;	2000US-0246523;
PR	08-NOV-2000;	2000US-0246524;
PR	08-NOV-2000;	2000US-0246525;
PR	08-NOV-2000;	2000US-0246526;
PR	08-NOV-2000;	2000US-0246527;
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PR	17-NOV-2000;	2000US-0249209;
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PR	17-NOV-2000;	2000US-0249211;
PR	17-NOV-2000;	2000US-0246611;
PR	08-NOV-2000;	2000US-0246613;
PR	17-NOV-2000;	2000US-0249212;
PR	17-NOV-2000;	2000US-0249213;
PR	17-NOV-2000;	2000US-0249214;
PR	17-NOV-2000;	2000US-0249215;
PR	17-NOV-2000;	2000US-0249216;
PR	17-NOV-2000;	2000US-0249217;
PR	17-NOV-2000;	2000US-0249218;
PR	17-NOV-2000;	2000US-0249219;
PR	17-NOV-2000;	2000US-0249220;
PR	17-NOV-2000;	2000US-0249221;
PR	17-NOV-2000;	2000US-0249222;
PR	17-NOV-2000;	2000US-0249223;
PR	17-NOV-2000;	2000US-0249224;
PR	17-NOV-2000;	2000US-0249225;
PR	17-NOV-2000;	2000US-0249226;
PR	17-NOV-2000;	2000US-0249227;
PR	17-NOV-2000;	2000US-0249228;
PR	17-NOV-2000;	2000US-0249229;
PR	17-NOV-2000;	2000US-0249230;

PR	17-NOV-2000;	2000US-0249300.
PR	01-DEC-2000;	2000US-0250160.
PR	01-DEC-2000;	2000US-0250391.
PR	05-DEC-2000;	2000US-0251030.
PR	05-DEC-2000;	2000US-0251988.
PR	05-DEC-2000;	2000US-0256719.
PR	06-DEC-2000;	2000US-0251479.
PR	08-DEC-2000;	2000US-0251856.
PR	08-DEC-2000;	2000US-0251868.
PR	08-DEC-2000;	2000US-0251869.
PR	08-DEC-2000;	2000US-0251989.
PR	08-DEC-2000;	2000US-0251990.
PR	11-DEC-2000;	2000US-0254097.
PR	05-JAN-2001;	2001US-0259678.
XX		
PA	(HUMA-)	HUMAN GENOME SCI INC.
XX		
XX	Rosen CA, Barash SC, Ruben	
PI	WPI; 2001-483426/52.	
XX		
XX	Nucleic acids encoding human	
PT	useful for preventing, diagnosing	
PT	metastasis -	
XX		
XX	Disclosure; SEQ ID NO 38705;	
XX	AAK54951 to AAK64702 encode t	
CC	amino acid sequences given in	
CC	activity, and can be used in	
CC	proteins and polynucleotides	
CC	treatment of diseases associated	
CC	example, they may be used to	
CC	expression by rectifying mutat	
CC	that affect the activity of (
CC	supplement the patients own p	
CC	polynucleotides may be used t	
CC	the nucleic acids into a host	
CC	protein. (I) proteins and pol	
CC	diagnose and treat immune/hae	
CC	cancers and cancer metastases	
CC	to AAK87694 represent human i	
CC	sequences from the present in	
CC	represent sequences used in t	
XX		
SQ	Sequence 3698 BP; 1067 A; 884	
	Query Match	17.0%;
	Best Local Similarity	55.6%;
	Matches	60; Conservative
QY	74 ATCTGGTGAGGCTATTATTATCCCC	
DB	590 AAGTAGTCTGCTGAATAAATACCC	
QY	134 CTGACTTTCATCTAGGAAGAAGCTA	
DB	650 CAAGTTCATTCTTAAGAAGACCT	
RESULT 14		
AAK64703		
ID	AAK64703 standard; DNA; 10548	
XX	AAK64703;	
XX		
AC		
DT	06-NOV-2001 (first entry)	
DE	Human immune/haematopoietic a	
XX		
KW	Human; immune; haematopoietic	
XX	cytostatic; gene therapy; vac	

OS Homo sapiens.
XX WO200157182-A2.
XX 09-AUG-2001.
XX 17-JAN-2001; 2001WO-US01354.
XX 31-JAN-2000; 2000US-0179065.
PR 04-FEB-2000; 2000US-0180628.
PR 24-FEB-2000; 2000US-0184664.
PR 02-MAR-2000; 2000US-0186350.
PR 16-MAR-2000; 2000US-0189874.
PR 17-MAR-2000; 2000US-0190076.
PR 18-APR-2000; 2000US-0198123.
PR 19-MAY-2000; 2000US-0205515.
PR 07-JUN-2000; 2000US-0209467.
PR 28-JUN-2000; 2000US-0214886.
PR 30-JUN-2000; 2000US-0215135.
PR 07-JUL-2000; 2000US-0216647.
PR 07-JUL-2000; 2000US-0216880.
PR 11-JUL-2000; 2000US-0217487.
PR 11-JUL-2000; 2000US-0217496.
PR 14-JUL-2000; 2000US-0218290.
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PR 26-JUL-2000; 2000US-0220964.
PR 14-AUG-2000; 2000US-0224518.
PR 14-AUG-2000; 2000US-0224519.
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PR 14-AUG-2000; 2000US-0225214.
PR 14-AUG-2000; 2000US-0225266.
PR 14-AUG-2000; 2000US-0225267.
PR 14-AUG-2000; 2000US-0225268.
PR 14-AUG-2000; 2000US-0225270.
PR 14-AUG-2000; 2000US-0225447.
PR 14-AUG-2000; 2000US-0225757.
PR 14-AUG-2000; 2000US-0225758.
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PR 08-SEP-2000; 2000US-0232081.
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PR 14-SEP-2000; 2000US-0232398.
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PR 14-SEP-2000; 2000US-0232400.
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PR 14-SEP-2000; 2000US-0233063.
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PR 21-SEP-2000; 2000US-0234223.
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PR 26-SEP-2000; 2000US-0235484.
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PR 29-SEP-2000; 2000US-0236368.
PR 29-SEP-2000; 2000US-0236369.
PR 29-SEP-2000; 2000US-0236370.
PR 02-OCT-2000; 2000US-0236802.
PR 02-OCT-2000; 2000US-0237037.
PR 02-OCT-2000; 2000US-0237038.
PR 02-OCT-2000; 2000US-0237039.
PR 02-OCT-2000; 2000US-0237040.
PR 13-OCT-2000; 2000US-0239935.
PR 13-OCT-2000; 2000US-0239937.
PR 20-OCT-2000; 2000US-0240960.
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PR 20-OCT-2000; 2000US-0241786.
PR 20-OCT-2000; 2000US-0241787.
PR 20-OCT-2000; 2000US-0241808.
PR 20-OCT-2000; 2000US-0241809.
PR 20-OCT-2000; 2000US-0241826.
PR 01-NOV-2000; 2000US-0244617.
PR 08-NOV-2000; 2000US-0246474.
PR 08-NOV-2000; 2000US-0246475.
PR 08-NOV-2000; 2000US-0246476.
PR 08-NOV-2000; 2000US-0246477.
PR 08-NOV-2000; 2000US-0246478.
PR 08-NOV-2000; 2000US-0246523.
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PR 08-NOV-2000; 2000US-0246532.
PR 08-NOV-2000; 2000US-0246609.
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PR 08-NOV-2000; 2000US-0246611.
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PR 17-NOV-2000; 2000US-0249207.
PR 17-NOV-2000; 2000US-0249208.
PR 17-NOV-2000; 2000US-0249209.
PR 17-NOV-2000; 2000US-0249210.
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PR 17-NOV-2000; 2000US-0249212.
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PR 17-NOV-2000; 2000US-0249214.
PR 17-NOV-2000; 2000US-0249215.
PR 17-NOV-2000; 2000US-0249216.
PR 17-NOV-2000; 2000US-0249217.
PR 17-NOV-2000; 2000US-0249218.
PR 17-NOV-2000; 2000US-0249244.
PR 17-NOV-2000; 2000US-0249245.
PR 17-NOV-2000; 2000US-0249264.
PR 17-NOV-2000; 2000US-0249265.
PR 17-NOV-2000; 2000US-0249297.
PR 17-NOV-2000; 2000US-0249299.
PR 17-NOV-2000; 2000US-0249300.
PR 01-DEC-2000; 2000US-0250160.
PR 01-DEC-2000; 2000US-0250391.
PR 05-DEC-2000; 2000US-0251030.
PR 05-DEC-2000; 2000US-0251988.
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PR 06-DEC-2000; 2000US-0251479.
PR 08-DEC-2000; 2000US-0251856.
PR 08-DEC-2000; 2000US-0251868.
PR 08-DEC-2000; 2000US-0251869.
PR 08-DEC-2000; 2000US-0251989.
PR 11-DEC-2000; 2000US-0251990.
PR 05-JAN-2001; 2000US-0254097.
PR 05-JAN-2001; 2000US-0259678.
PR (HUMA-) HUMAN GENOME SCI INC.
XX PA

PR 20-OCT-2000; 2000US-0241785.
PR 20-OCT-2000; 2000US-0241786.
PR 20-OCT-2000; 2000US-0241787.
PR 20-OCT-2000; 2000US-0241808.
PR 20-OCT-2000; 2000US-0241809.
PR 20-OCT-2000; 2000US-0241826.
PR 01-NOV-2000; 2000US-0244617.
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PR 08-NOV-2000; 2000US-0246476.
PR 08-NOV-2000; 2000US-0246477.
PR 08-NOV-2000; 2000US-0246478.
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PR 08-NOV-2000; 2000US-0246611.
PR 08-NOV-2000; 2000US-0246613.
PR 17-NOV-2000; 2000US-0249207.
PR 17-NOV-2000; 2000US-0249208.
PR 17-NOV-2000; 2000US-0249209.
PR 17-NOV-2000; 2000US-0249210.
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PR 17-NOV-2000; 2000US-0249214.
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PR 17-NOV-2000; 2000US-0249217.
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PR 17-NOV-2000; 2000US-0249244.
PR 17-NOV-2000; 2000US-0249245.
PR 17-NOV-2000; 2000US-0249264.
PR 17-NOV-2000; 2000US-0249265.
PR 17-NOV-2000; 2000US-0249297.
PR 17-NOV-2000; 2000US-0249299.
PR 01-DEC-2000; 2000US-0250160.
PR 01-DEC-2000; 2000US-0250391.
PR 05-DEC-2000; 2000US-0251030.
PR 05-DEC-2000; 2000US-0251988.
PR 06-DEC-2000; 2000US-0251479.
PR 08-DEC-2000; 2000US-0251856.
PR 08-DEC-2000; 2000US-0251868.
PR 08-DEC-2000; 2000US-0251869.
PR 08-DEC-2000; 2000US-0251989.
PR 08-DEC-2000; 2000US-0251990.
PR 11-DEC-2000; 2000US-0254097.
PR 05-JAN-2001; 2001US-0259678.
(HUMA-) HUMAN GENOME SCI INC.

Rosen CA, Barash SC, Ruben SM;
WPI; 2001-483426/52.

Nucleic acids encoding human immune/hematopoietic antigen polypeptides,
useful for preventing, diagnosing and/or treating cancers and
metastasis -

Disclosure; SEQ ID NO 24625; 3071pp + Sequence Listing; English.

AAK54951 to AAK64702 encode the human immune/hematopoietic antigen (I)
amino acid sequences given in AAK62170 to AAK61921. (I) have cytostatic
activity, and can be used in gene therapy and vaccine production. (I)
proteins, and polynucleotides may be used in the prevention, diagnosis and
treatment of diseases associated with inappropriate (I) expression. For

CC example, they may be used to treat disorders associated with decreased
CC expression by rectifying mutations or deletions in a patient's genome
CC that affect the activity of (I) by expressing inactive proteins or to
CC supplement the patients own production of (I). Additionally, (I)
CC polynucleotides may be used to produce the secreted (I) by inserting the
CC the nucleic acids into a host cell and culturing the cell to express the
CC protein. (I) proteins and polynucleotides may be used to prevent,
CC diagnose and treat immune/hematopoietic-related diseases, especially
CC cancers and cancer metastases of haematopoietic-derived cells. AAK64703
CC to AAK87694 represent human immune/haematopoietic antigen genomic
CC sequences from the present invention. AAK54942 to AAK54950 and AAK62169
CC represent sequences used in the exemplification of the present invention.
XX

Sequence 10548 BP; 3143 A; 2302 C; 2158 G; 2945 T; 0 other;

Query Match 17.0%; Score 31.2; DB 22; Length 10548;
Best Local Similarity 55.6%; Pred. No. 3.9;
Matches 60; Conservative 0; Mismatches 48; Indels 0; Gaps 0;
QY 74 ATGTGGTGAGGCTATTATTTCCCTCCAGTGGCTTCCCTGGGCTATGGATGAACAGTGG 133
Db 7440 AAGTAGTTCTGCTAATAATCACCCTAGTCCCTGATTGTTATGAGCAGGCGAAGCTGTAG 7499
QY 134 CTGACTTCATCTAGGAAGAGCTATGGCTTCTGCTCTCCTGGAGCTCAC 181
Db 7500 CAAAGTTCTATTCTAAAGAACCTAAATGGCGCGTGTGTGGTGGCTCAC 7547

Search completed: May 25, 2003, 03:13:59
Job time : 250 secs

GenCore version 5.1.4_p5_4578
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: May 25, 2003, 03:09:12 ; Search time 1754 Seconds
(without alignments)
1689.723 Million cell updates/sec

Title: us-09-660-568-49

Perfect score: 183

Sequence: 1 cacacactccccattctga.....ctgtctctcggagctcaaca 183

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

- EST:*
- 1: em_estba:*
 - 2: em_esthum:*
 - 3: em_estin:*
 - 4: em_estmu:*
 - 5: em_estov:*
 - 6: em_estpl:*
 - 7: em_estro:*
 - 8: em_hic:*
 - 9: gb_est1:*
 - 10: gb_est2:*
 - 11: gb_hic:*
 - 12: gb_est3:*
 - 13: gb_est4:*
 - 14: gb_est5:*
 - 15: em_estfun:*
 - 16: em_estom:*
 - 17: gb_gss:*
 - 18: em_gss_hum:*
 - 19: em_gss_inv:*
 - 20: em_gss_pin:*
 - 21: em_gss_vrt:*
 - 22: em_gss_fun:*
 - 23: em_gss_mam:*
 - 24: em_gss_mus:*
 - 25: em_gss_other:*
 - 26: em_gss_pro:*
 - 27: em_gss_rod:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
c 1	41.2	22.5	961	12	BG775699 602650070
c 2	37.6	20.5	501	17	BH309749 CH230-181
c 3	37	20.2	506	17	AQ419671 RPCI-11-1
c 4	36.6	20.0	789	17	BH317352 CH230-119
c 5	35.4	19.3	702	12	BG719810 6026501217
c 6	34.2	18.7	831	12	BF571585 602078004

7	34	18.6	283	12	BG161919
8	33.6	18.4	695	9	AI512580
c 9	33	18.0	438	9	AI243308
c 10	33	18.0	467	9	AI806410
c 11	33	18.0	481	17	BH274856
c 12	33	18.0	594	17	BH268816
c 13	32.8	17.9	582	13	BM663839
c 14	32.8	17.9	994	17	CNS0174A
c 15	32.4	17.7	226	17	AQ543989
c 16	32.4	17.7	524	17	B01998
c 17	32.4	17.7	544	17	AQ809155
c 18	32.4	17.7	609	12	BG720090
c 19	32	17.5	365	17	AZ745700
c 20	32	17.5	401	17	BH305232
c 21	31.8	17.4	455	17	AQ536443
c 22	31.8	17.4	591	12	BG606222
c 23	31.6	17.3	638	17	AG034524
c 24	31.6	17.3	649	13	BM416296
c 25	31.6	17.3	720	17	BH307258
c 26	31.4	17.2	258	14	H32274
c 27	31.4	17.2	308	17	AQ252041
c 28	31.4	17.2	574	13	BI359567
c 29	31.4	17.2	623	10	BE082216
c 30	31.4	17.2	634	9	AL510875
c 31	31.4	17.2	695	17	BH364239
c 32	31.2	17.0	316	17	AQ584142
c 33	31.2	17.0	446	17	AQ977475
c 34	31.2	17.0	524	17	AQ177467
c 35	31.2	17.0	535	17	AQ284523
c 36	31.2	17.0	751	17	AZ633721
c 37	31	16.9	502	17	AZ217607
c 38	31	16.9	566	17	AZ792647
c 39	31	16.9	1082	17	CNS078VY
c 40	30.8	16.8	417	12	BG190341
c 41	30.8	16.8	425	17	BH258367
c 42	30.8	16.8	481	17	BH860938
c 43	30.8	16.8	514	17	AQ792988
c 44	30.8	16.8	517	14	BQ090484
c 45	30.8	16.8	577	17	AQ189764

ALIGNMENTS

RESULT 1
BG775699/c
LOCUS 602650070T1 NIH_MGC_40 Homo sapiens cDNA clone IMAGE:4760993 3',
DEFINITION BG775699 961 bp mRNA linear EST 15-MAY-2001
ACCESSION BG775699
VERSION BG775699.1 GI:14046016.
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 961)
AUTHORS NIH-MGC http://mgc.nci.nih.gov/.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cygabs-remail.nih.gov
Tissue Procurement: DCTD/DTP
CDNA Library Preparation: Ling Hong/Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LICM1612 row: 1 column: 18
High quality sequence start: 4
High quality sequence stop: 783.
Location/Qualifiers

FEATURES

Db 645 CTTTTCCTGCTGCAAGAGACCTGCCCTGGTGAACTCGGGACACACAGAGGAGAAATTCCTC 586

BF571585	LOCUS	DEFINITION
----------	-------	------------

BF571585	LOCUS	BF571585	831 bp	mrna	linear	EST 12-DEC-2000
DEFINITION	602078004F1	NIH MGC 62	Homo sapiens	CDNA clone	IMAGE:4252465	5'

```

mRNA sequence.
ACCESSION BF571585
VERSION BF571585.1 GI:11645297
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
TITLE NIH-MGC http://mgi.nci.nih.gov/.
JOURNAL 1 (bases 1 to 831)
COMMENT National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Tissue Procurement: ATCC/DCTD/DTF
Email: cgapbs-re@mail.nih.gov
CDNA Library Preparation: CLONETECH Laboratories, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLCM1077 row: h column: 02
High quality sequence stop: 571..
FEATURES
Location/Qualifiers
1..831
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="IMAGE:4252465"
/clone_lib="NIH-MGC-62"
/tissue_type="melanotic melanoma, high MDR"
/lab_host="DH10B (T1 phage-resistant)"
/Note="Organ: skin; Vector: pDMR-LIB (Clontech); Site:1:
SfiI (ggccgctggcc); Site:2: SfiI (ggccattatggcc);
Double-stranded cDNA was prepared from cell line RNA.
5' and 3' adaptors were used in cloning as follows: 5'
adaptor sequence: 5'-CACGGCCATTATGGCC-3' and 3' adaptor
sequence: 5'-ATTCCTAGAGCGGCGGCGGACATG-dT(30)BN-3'
(where B = A, C, G, or T). Average
insert size 1.75 kb (range 0.9-4.0 kb). 15/15 colonies
contained inserts by PCR. This library was enriched for
full-length clones and was constructed by Clontech
Laboratories (Palo Alto, CA)."
BASE COUNT 219 a 190 c 216 g 206 t
ORIGIN
Query Match 18.7%; Score 34.2; DB 12; Length 831;
Best Local Similarity 58.3%; Pred. No. 7.9;
Matches 60; Conservative 0; Mismatches 43; Indels 0; Gaps 0;

64 CAGAAGGAGATGTGTGGAGGCTATTATTCCTCCAGTCCCTGCTGGCTATGGA 123
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 334 CAGATCGAGGACTGTGTGGTGTATTATTCCTCCATTGGCTAAACAGATGAACAGCGA 393

QY 124 TGAACAGTGCACATTCATAGAAAGAGCTATGGCTTCG 166
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 394 GGTGTCTACACTAACAGCAGCAGCTGGGAAGAGATTAGCTGTG 436

RESULT 7
BG161919
LOCUS BG161919 283 bp mRNA linear EST 06-FEB-2001
DEFINITION dc43b12.y1 NICHD XGC Emb3 Xenopus laevis cDNA clone IMAGE:3399886
5', mRNA sequence.
ACCESSION BG161919
VERSION BG161919.1 GI:12695838
KEYWORDS EST.
SOURCE African clawed frog.
ORGANISM Xenopus laevis
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae;
Xenopodinae; Xenopus.
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
Other ESTs: dc43b12.x1
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-re@mail.nih.gov
Tissue Procurement: Martha Rebert, Steven L. Klein, Ph.D.
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: Xenopus clones from this library are available
through the I.M.A.G.E. Consortium/LLNL at: info@image.llnl.gov
Seq primer: -40RP from Gibco
High quality sequence stop: 269.
FEATURES
Location/Qualifiers
1..283
/organism="Xenopus laevis"
/db_xref="taxon:8355"
/clone_lib="IMAGE:3399886"
/clone_lib="NICHD XGC Emb3"
/tissue_type="embryo (stages 24-25)"
/lab_host="DH10B (phage-resistant)"
/Note="Vector: pCMV-SPORT6; Site:1: NotI; Site:2: SalI;
Cloned unidirectionally. Primer: Oligo dT. Average insert
size 1.7 kb. Constructed by Life Technologies. Note: This
is a Xenopus Gene Collection (XGC) library."
BASE COUNT 76 a 58 c 68 g 81 t
ORIGIN
Query Match 18.6%; Score 34; DB 12; Length 283;
Best Local Similarity 61.1%; Pred. No. 5.1;
Matches 55; Conservative 0; Mismatches 35; Indels 0; Gaps 0;

QY 63 GCAGAGGAGATGTGTGGAGGCTATTATTCCTCCAGTCCCTGCTGGCTATGG 122
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 157 GCAGATGAAGATTGTGAAGGCTATTATTCACAGAGCAATGCTCTCTGGCTTCAT 216

QY 123 ATGACAGTGGCTGACTTCATCTAGGAAG 152
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 217 ATTGTAGAGGCTTGTAGACCAAGGAAG 246

RESULT 8
AI512580
LOCUS AI512580 695 bp mRNA linear EST 23-APR-2001
DEFINITION LD44491.Sprime LD Drosophila melanogaster embryo pot2 Drosophila
melanogaster cDNA clone LD44491 5 similar to Cyp310a1: FBan0010391
'cytochrome P450' located on: 2L 37A3-37A3:: 04/10/2001, mRNA
sequence.
ACCESSION AI512580
VERSION AI512580.2 GI:13770009
KEYWORDS EST.
SOURCE fruit fly.
ORGANISM Drosophila melanogaster
REFERENCE Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
AUTHORS Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
Ephydroidea; Drosophilidae; Drosophila.
Lewis, S. and Rubin, G.M.
Harvey, D., Brokstein, P., Hong, L., Evans-Holm, M., Su, C., Tsang, G.,
BDGP/HMI Drosophila EST Project
Unpublished (2001)
On Mar 16, 1999 this sequence version replaced gi:4421998.
Other ESTs: LD44491.Sprime
Contact: Stapleton, M.
BDGP
Lawrence Berkeley National Lab
One Cyclotron Rd, Berkeley, CA 94720, USA
Fax: 510 486 6798
Email: http://www.fruitfly.org/EST, est@fruitfly.berkeley.edu
hit genomic AE003659: arm:2L [18246373,18507269]
estimated-cyto:36F7-37A4: 04/10/2001
Plate: LD.444 row: H column: 7

```


High quality sequence stop: 585

FEATURES
source

Location/Qualifiers

1. .695
/organism="Drosophila melanogaster"
/db_xref="taxon:7227"
/clone="LD4491"
/clone_lib="LD Drosophila melanogaster embryo pot2"
/sex="male and female"
/dev_stage="0 to 24 hours mixed stage embryonic"
/lab_host="XLI Blue"
/note="Organ: Embryo; Vector: pOT2; Site:1: EcoRI; Site:2:
XhoI; Sized fractionated cDNAs were directly ligated into
pOT2."

BASE COUNT 171 a 175 c 176 g 173 t

ORIGIN

Query Match 18.4%; Score 33.6; DB 9; Length 695;

Best Local Similarity 53.9%; Pred. No. 11;
Matches 69; Conservative 0; Mismatches 59; Indels 0; Gaps 0;

38 CCTAAGATGTCAGAGATCCAAAGTCAGAGAGAGAGATGTGGTGAGGCTATTATTCGCC 97

Db 400 CCAACAGATTGTCAGAGATGTCGGCAGGTCAGAGAGAGAGATGTGGTGAGGCTATTATTCACC 459

Qy 98 CAGTGCCTTCCTGCTGGCTATGATGACAGTGGCTGACTTCATCTAGGAAAGAGCTA 157

Db 460 ATTTGCTGGACAGGCAAGCGCATGATCCACAGGAAGTGGATATCCAGCAAAACTCA 519

Qy 158 TGGCTTCT 165

Db 520 GAGTTTAT 527

RESULT 9

AI243308/c

LOCUS

DEFINITION q41b03.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone

IMAGE:1847213 3', mRNA sequence.

AI243308

AI243308.1 GI:3838705

VERSION

KEYWORDS

SOURCE

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Tumor Gene Index

Unpublished (1997)

Contact: Robert Strausberg, Ph.D.

Email: cgapbs-r@mail.nih.gov

This clone is available royalty-free through LLNL; contact the

IMAGE Consortium (info@image.llnl.gov) for further information.

Insert Length: 623 Std Error: 0.00

Seq primer: -40UP from Gibco

High quality sequence stop: 415.

Location/Qualifiers

1. .438

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:1847213"

/clone_lib="Soares_NFL_T_GBC_S1"

/lab_host="DH10B"

/note="Organ: pooled; Vector: pT7T3D-Pac (Pharmacia) with

a modified polylinker; Site:1: Not I; Site:2: Eco RI;

Equal amounts of plasmid DNA from three normalized

libraries (fetal lung NDHL19W, testis NHT, and B-cell

NCI-CGAP-GB1) were mixed, and ss circles were made in

vitro. Following HAP purification, this DNA was used as

tracer in a subtractive hybridization reaction. The driver

was PCR-amplified cDNAs from pools of 5,000 clones made

from the same 3 libraries. The pools consisted of
I.M.A.G.E. clones 297480-302087, 682632-687239,
726408-728711, and 729096-731399. Subtraction by Bento
Soares and M. Fatima Bonaldo.

BASE COUNT 156 a 85 c 78 g 119 t

ORIGIN

Query Match 18.0%; Score 33; DB 9; Length 438;

Best Local Similarity 57.1%; Pred. No. 13;

Matches 60; Conservative 0; Mismatches 45; Indels 0; Gaps 0;

Qy 41 AAGATGTCCAGAGATCCAAAGTCAGAGAGAGAGATGTGGTGAGGCTATTATTCGCCAG 100

Db 202 AAGTACTAGTGTATAGAGATGGGGAGAGAGATAAGTTAAATCATTTATTCCTCGG 143

Qy 101 TGCCTTCCTGCTGGCTATGGATGACAGTGGCTGACTTCATCT 145

Db 142 CTCCTTCCTTCTGGCTATGGCTGAGTGGTGGTTCCTTTCT 98

RESULT 10

AI806410/c

LOCUS

DEFINITION wf27f08.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone

IMAGE:2356839 3', mRNA sequence.

AI806410

AI806410.1 GI:5392976

VERSION

KEYWORDS

SOURCE

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Tumor Gene Index

Unpublished (1997)

Contact: Robert Strausberg, Ph.D.

Email: cgapbs-r@mail.nih.gov

This clone is available royalty-free through LLNL; contact the

IMAGE Consortium (info@image.llnl.gov) for further information.

Insert Length: 619 Std Error: 0.00

Seq primer: -40UP from Gibco

High quality sequence stop: 415.

Location/Qualifiers

1. .467

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:2356839"

/clone_lib="Soares_NFL_T_GBC_S1"

/lab_host="DH10B"

/note="Organ: pooled; Vector: pT7T3D-Pac (Pharmacia) with

a modified polylinker; Site:1: Not I; Site:2: Eco RI;

Equal amounts of plasmid DNA from three normalized

libraries (fetal lung NDHL19W, testis NHT, and B-cell

NCI-CGAP-GB1) were mixed, and ss circles were made in

vitro. Following HAP purification, this DNA was used as

tracer in a subtractive hybridization reaction. The driver

was PCR-amplified cDNAs from pools of 5,000 clones made

I.M.A.G.E. clones 297480-302087, 682632-687239,

726408-728711, and 729096-731399. Subtraction by Bento

Soares and M. Fatima Bonaldo.

BASE COUNT 167 a 88 c 81 g 131 t

ORIGIN

Query Match 18.0%; Score 33; DB 9; Length 467;

Best Local Similarity 57.1%; Pred. No. 14;

Matches 60; Conservative 0; Mismatches 45; Indels 0; Gaps 0;

Qy 41 AAGATGTCCAGAGATCCAAAGTCAGAGAGAGAGATGTGGTGAGGCTATTATTCGCCAG 100

Db 203 AAGTACTAGTGTATAGAGATGGGGAGAGAGATAAGTTAAATCATTTATTCCTCGG 144

AUTHORS TITLE JOURNAL MEDLINE COMMENT

Bonaldo, M.F., Lennon, G. and Soares, M.B.
Normalization and subtraction: two approaches to facilitate gene
discovery
Genome Res. 6 (9), 791-806 (1996)
97044477
Contact: Soares, MB
Program for Rat Gene Discovery and Mapping
University of Iowa
451 Eckstein Medical Research Building Iowa City, IA 52242, USA
Tel: 319 335 8250
Fax: 319 335 9565
Email: msoares@blue.weeg.uiowa.edu
Tissue Procurement: Dr. Gregg Hagaman
cDNA Library Preparation: Dr. M. Bento Soares, University of Iowa
DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
Clone Distribution: Dr. M. Bento Soares, University of Iowa
Genetics (www.resgen.com).
The following repetitive elements were found in this cDNA
sequence: 11-93, >ALU (matched complement)
Seq primer: M13 Forward
POLYA-Yes.

FEATURES source

Location/Qualifiers
1. .582
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="UI-E-CL1-afe-k-03-0-UI"
/clone_lib="UI-E-CL1"
/tissue_type="human retina"
/dev_stage="adult"
/lab_host="DH10B (Life Technologies) (T1 phage resistant)"
/note="Organ: eye; Vector: pT73-Pac (Pharmacia) with a
modified polylinker; Site_1: EcoR I; Site_2: Not I;
UI-E-CL1 is a normalized cDNA library containing the
following tissue(s): retina. The library was constructed
according to Bonaldo, Lennon and Soares, Genome Research,
6:791-806, 1996. First strand cDNA synthesis was primed
with an oligo-dT primer containing a Not I site. Double
stranded cDNA was ligated to an EcoR I adaptor, digested
with Not I, and cloned directionally into pT73-Pac
vector. The oligonucleotide used to prime the synthesis of
first-strand cDNA contains a library tag sequence that is
located between the Not I site and the (dT)18 tail. The
sequence tag for this library is CCGCG. This library was
created for the program, Gene Discovery in the Visual
System, supported by National Eye Institute (NEI).
TAG LIB=UI-E-CL1
TAG TISSUE=human retina
TAG_SEQ=CCGCG"

BASE COUNT 171 a 128 c 98 g 185 t
ORIGIN

Query Match 17.9%; Score 32.8; DB 13; Length 582;
Best Local Similarity 54.0%; Pred. No. 18;
Matches 67; Conservative 0; Mismatches 57; Indels 0; Gaps 0;
Qy 8 TCCCCCATCTGAGCCCAAGAGCGTCATCCCTTAAGAGATGTCACAGATCCAAAGTCAGA 67
Db 149 TCCTTCATCAAGCGACAGCGTGGTTCATTAGTAAGCTTTGGCATACTGCTAATGGCA 90
Qy 68 AGGAGATGGTGGAGGCTATTTATCCCCCAGTCGCTTCCTCTGGCTATGGATGAA 127
Db 89 GGCAGAGTTTGCAGTGAGTCAGATTCGCCACTGCACTCCAGCTGGCGACTGAGCAA 30
Qy 128 CAGT 131
Db 29 GACT 26

RESULT 14 CNS0174A/c LOCUS DEFINITION

CNS0174A 994 bp DNA linear GSS 26-JUL-1999
Drosophila melanogaster genome survey sequence SP6 end of BAC

ACCESSION VERSION KEYWORDS SOURCE ORGANISM

AL107572
AL107572.1 GI:5627876
GSS.
Drosophila melanogaster.
Drosophila melanogaster.

REFERENCE AUTHORS TITLE JOURNAL

Submitted (23-JUL-1999) Genoscope - Centre National de Sequencage :
BP 191 91006 Evry cedex - FRANCE (E-mail : seqref@genoscope.cns.fr)
- Web : www.genoscope.cns.fr
Determination of this BAC-end sequence was carried out as part of a
collaboration with the European Drosophila Genome Project (EDGP) -
http://www.edgp.ebi.ac.uk -. This Drosophila melanogaster BAC
library (Dros BAC) was made by Alain Billaud at CEPH (Centre
d'Etude du Polymorphisme Humain) with funding provided by a MRC
project grant. The DNA was prepared from embryos by Alain Bucheton
and Genevieve Payan. It has been constructed in the vector
pBelobAC11.

FEATURES source

Location/Qualifiers
1. .994
/organism="Drosophila melanogaster"
/db_xref="taxon:7227"
/clone="BACN17G20"
/clone_lib="DrosBAC"
/plasmid="pBelobAC11"
/notes="end : SP6"

BASE COUNT 223 a 214 c 242 g 250 t 65 others
ORIGIN

Query Match 17.9%; Score 32.8; DB 17; Length 994;
Best Local Similarity 55.2%; Pred. No. 24;
Matches 64; Conservative 0; Mismatches 52; Indels 0; Gaps 0;
Qy 38 CCTAAGATGTCACAGATCCCAAGTCGAGAGGAATGTGTGAGGCTATTATTCCCC 97
Db 427 CCACAGATGTCGAGATGTCGGCAGGTCACAGAAAGTGGTCATTAATTCACC 368
Qy 98 CAGTGCCTTCCTGCTGGCTATGATGAACAGTGGCTGACTATCTAGGAAGA 153
Db 367 ATTGCTGGACAGGCAAGGCCAAGGATGTCCACAGGAAGTGATCCAGCAAAA 312

RESULT 15 AQ543989/c LOCUS

DEFINITION
RPCI-11-357B13-TV RPCI-11 Homo sapiens genomic clone RPCI-11-357B13
DNA sequence.
ACCESSION
AQ543989
KEYWORDS
GSS.
SOURCE
human.
ORGANISM
Homo sapiens

REFERENCE AUTHORS TITLE JOURNAL COMMENT

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 226)
Zhao, S., Adams, M.D., Nierman, W., Malek, J., de Jong, P. and Venter
J.C.
Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready
Map Building
Unpublished (1997)
Other_GSSs: RPCI-11-357B13.TJ
Contact: Shaying Zhao, William Nierman, Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850
Tel: 301 838 0200
Fax: 301 838 0208

GenCore version 5.1.4_p5.4578
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OM nucleic - nucleic search, using sw model

Run on: May 25, 2003, 03:09:47 ; Search time 52 Seconds
(without alignments)
1079.266 Million cell updates/sec

Title: US-09-660-568-49

Perfect score: 183

Sequence: 1 cacacactccccattctga.....ctgtctctctggagctcacca 183

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 441362 seqs, 153338381 residues

Total number of hits satisfying chosen parameters: 882724

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents NA:*

1: /cgn2_6/ptodata/1/ina/5A_COMB.seq:*

2: /cgn2_6/ptodata/1/ina/5B_COMB.seq:*

3: /cgn2_6/ptodata/1/ina/6A_COMB.seq:*

4: /cgn2_6/ptodata/1/ina/6B_COMB.seq:*

5: /cgn2_6/ptodata/1/ina/PCTUS_COMB.seq:*

6: /cgn2_6/ptodata/1/ina/backfiles1.seq:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	183	100.0	183	4	US-09-046-894-49
2	28.2	15.4	2846	4	Sequence 49, Appl
3	28	15.3	2297	4	Sequence 1, Appl
4	27.6	15.1	2367	3	Sequence 12, Appl
5	27.4	15.0	3243	2	Sequence 17, Appl
6	27.4	15.0	3243	2	Sequence 6, Appl
7	27.4	15.0	3243	3	Sequence 6, Appl
8	27	14.8	246240	2	Sequence 6, Appl
9	27	14.8	246240	2	Sequence 20, Appl
10	27	14.8	246240	2	Sequence 21, Appl
11	26.6	14.5	2624	1	Sequence 22, Appl
12	26.6	14.5	2624	3	Sequence 15, Appl
13	26	14.2	3138	4	Sequence 15, Appl
14	26	14.2	3834	3	Sequence 1, Appl
15	26	14.2	3854	1	Sequence 18, Appl
16	26	14.2	3858	2	Sequence 1, Appl
17	26	14.2	3858	4	Sequence 98, Appl
18	26	14.2	3863	4	Sequence 8, Appl
19	26	14.2	3863	6	Sequence 1, Appl
20	25.8	14.1	2380	6	Patent No. 5217870
21	25.6	14.0	1473	4	Patent No. 5268463
22	25.6	13.9	1001	4	Sequence 7, Appl
23	25.4	13.9	1001	4	Sequence 214, Appl
24	25.4	13.9	2777	4	Sequence 215, Appl
25	25.4	13.9	2777	4	Sequence 3, Appl
26	25.4	13.9	2777	4	Sequence 310-463-3
27	25.4	13.9	2790	3	Sequence 2, Appl
28	25.4	13.9	2922	4	Sequence 21, Appl
29	25.4	13.9	2922	4	Sequence 1, Appl

c 28	25.4	13.9	2922	4	US-08-842-248A-1	Sequence 1, Appl
29	25.4	13.9	111282	4	US-09-754-250-3	Sequence 3, Appl
c 30	25.2	13.8	285	4	US-09-134-001C-2573	Sequence 2573, Ap
c 31	25.2	13.8	441	4	US-09-134-001C-2432	Sequence 2432, Ap
c 32	25.2	13.8	1001	4	US-09-641-638-250	Sequence 250, App
c 33	25.2	13.8	1001	4	US-09-641-638-258	Sequence 258, App
c 34	25.2	13.8	1512	3	US-08-803-603-4	Sequence 4, Appl
c 35	25.2	13.8	1515	3	US-08-369-822C-7	Sequence 7, Appl
c 36	25.2	13.8	1515	3	US-08-582-776C-7	Sequence 7, Appl
c 37	25.2	13.8	1515	3	US-08-434-831B-7	Sequence 7, Appl
c 38	25.2	13.8	1521	3	US-08-779-764A-10	Sequence 10, Appl
c 39	25.2	13.8	1521	3	US-08-779-764A-11	Sequence 11, Appl
c 40	25.2	13.8	1521	3	US-08-779-764A-12	Sequence 12, Appl
c 41	25.2	13.8	8910	3	US-08-369-822C-19	Sequence 19, Appl
c 42	25.2	13.8	8910	3	US-08-582-776C-19	Sequence 19, Appl
c 43	25.2	13.8	8910	3	US-08-434-831B-19	Sequence 19, Appl
c 44	25.2	13.8	9126	1	US-08-580-038-26	Sequence 26, Appl
45	25.2	13.8	9126	2	US-08-639-857-3	Sequence 3, Appl

ALIGNMENTS

RESULT 1

US-09-046-894-49

; Sequence 49, Application US/09046894

; Patent No. 6190857

; GENERAL INFORMATION:

; APPLICANT: Ralph, David

; APPLICANT: An' Gang

; APPLICANT: O'Hara, Mark S.

; APPLICANT: Veltri, Robert

; TITLE OF INVENTION: DIAGNOSIS OF DISEASE STATE USING mRNA

; NUMBER OF SEQUENCES: 55

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Arnold, White & Durkee

; STREET: P.O. Box 4433

; CITY: Houston

; STATE: Texas

; COUNTRY: USA

; ZIP: 77210

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: PatentIn Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/09/046,894

; FILING DATE: Concurrently Herewith

; CLASSIFICATION:

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: US 60/041,576

; FILING DATE: 24-MAR-1997

; ATTORNEY/AGENT INFORMATION:

; NAME: Nakashima, Richard A.

; REGISTRATION NUMBER: P-42,023

; REFERENCE/DOCKET NUMBER: UROC:014

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (512) 418-3000

; TELEFAX: (512) 474-7577

; INFORMATION FOR SEQ ID NO: 49:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 183 base pairs

; TYPE: nucleic acid

; STRANDEDNESS: single

; TOPOLOGY: linear

; US-09-046-894-49

Query Match 100.0%; Score 183; DB 4; Length 183;

Best Local Similarity 100.0%; Pred. No. 8.6e-56;

Matches 183; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

[illegible]

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/276,968A
FILING DATE: 19-JUL-1994
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/905,795
FILING DATE: 29-JUN-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/016,863
FILING DATE: 10-FEB-1993
ATTORNEY/AGENT INFORMATION:

```

; ATTORNEY/AGENT INFORMATION:
; NAME: Kurt G. Briscoe
; REGISTRATION NUMBER: 33,141
; REFERENCE/DOCKET NUMBER: MDI 251.4-KGB
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (914) 332-1700
; TELEFAX: (914) 332-1844
; INFORMATION FOR SEQ ID NO: 6:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 3243 nucleotides
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; US-08-239-276-6

Query Match 15.0%; Score 27.4; DB 2; Length 3243;
Best Local Similarity 57.6%; Pred.No.5.4;
Matches 49; Conservative 0; Mismatches 36; Indels 0; Gaps

QY 35 ATCCCTAAGGATGTCCAGAGATCCAAGTGCAGAGAAGGAGAAATGTGGTGAGCGCTATTATTTC 94
   ||| |||| | |||| | |||| | |||| | |||| | |||| | |||| | |||| |
Db 161 ATGTCAAAGGACCAAGAGAGCCTAGCAGAGAAGGAGCTCCAGCTTCTGGTCATGATTC 220

QY 95 CCCAGTGCCTTCCTGCTGGGGCTA 119
   || | | | | | | | | | | |
Db 221 ACCAGCTGTCCACCCTCGGGACCA 245

RESULT 6
US-08-468-579B-6
; Sequence 6, Application US/08468579B
; Patent No. 5981700
; GENERAL INFORMATION:
; APPLICANT: Rabin, Daniel
; TITLE OF INVENTION: PANCREATIC ISLET CELL ANTIGENS
; TITLE OF INVENTION: OBTAINED BY MOLECULAR CLONING
; NUMBER OF SEQUENCES: 19
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Sprung Kramer Schaefer & Briscoe
; STREET: 660 White Plains Road
; CITY: Tarrytown
; STATE: New York
; COUNTRY: USA
; ZIP: 10591-5144
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.50 inch, 1.4 Mb storage.
; COMPUTER: Apple Macintosh
; OPERATING SYSTEM: System 7.5
; SOFTWARE: WordPerfect
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/468,579B
; FILING DATE: 06-JUN-1995
; CLASSIFICATION: 530
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/239,276
; FILING DATE: 05-MAY-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/715,181
; FILING DATE: 14-JUN-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/441,703
; FILING DATE: 04-DEC-1989
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/312,543
; FILING DATE: 17-FEB-1989
; ATTORNEY/AGENT INFORMATION:
; NAME: Kurt G. Briscoe
; REGISTRATION NUMBER: 33,141
; REFERENCE/DOCKET NUMBER: MDI 251.5-KGB
; TELECOMMUNICATION INFORMATION:

```

TELEPHONE: (914) 332-1700
TELEFAX: (914) 332-1844
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 3243 nucleotides
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-468-579B-6

Query Match 15.0%; Score 27.4; DB 2; Length 3243;
Best Local Similarity 57.6%; Pred. No. 5.4; Mismatches 0; Gaps 0;
Matches 49; Conservative 0;

QY 35 ATCCCTAAGGATGTCAGAGATCCCAAGTGCAGAGGAGAATGTGTGAGGCTATTATTC 94
DB 161 ATGTCAAGGGACCAAGAGACCCCTAGCAGAGAGAGAGCTCCAGCTTCGTGTCATGATTC 220
QY 95 CCCAGTGCCTTCCTGCTGGGCTA 119
DB 221 ACCAGCTGTCCACCTCGGGACCA 245

RESULT 7

US-08-468-577B-6
Sequence 6, Application US/08468577B
Patent No. 6001804

GENERAL INFORMATION:
APPLICANT: Rabid, Daniel
TITLE OF INVENTION: PANCREATIC ISLET CELL ANTIGENS
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESS: Sprung Kramer Schaefer & Briscoe
STREET: 660 White Plains Road
CITY: Tarrytown
STATE: New York
COUNTRY: USA
ZIP: 10591-5144

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.50 inch, 1.4 Mb storage
COMPUTER: Apple Macintosh
OPERATING SYSTEM: System 7.5
SOFTWARE: WordPerfect
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/468,577B
FILING DATE: 06-JUN-1995
CLASSIFICATION: 514

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/239,276
FILING DATE: 03-MAY-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/872,646
FILING DATE: 08-JUN-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/715,181
FILING DATE: 14-JUN-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/441,703
FILING DATE: 04-DEC-1989
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/312,543
FILING DATE: 17-FEB-1989

ATTORNEY/AGENT INFORMATION:
NAME: Kurt G. Briscoe
REGISTRATION NUMBER: 33,141
REFERENCE/DOCKET NUMBER: MDI 251.8-KGB
TELECOMMUNICATION INFORMATION:
TELEPHONE: (914) 332-1700
TELEFAX: (914) 332-1844
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 3243 nucleotides

TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-468-577B-6

Query Match 15.0%; Score 27.4; DB 3; Length 3243;
Best Local Similarity 57.6%; Pred. No. 5.4; Mismatches 0; Gaps 0;
Matches 49; Conservative 0;

QY 35 ATCCCTAAGGATGTCAGAGATCCCAAGTGCAGAGGAGAATGTGTGAGGCTATTATTC 94
DB 161 ATGTCAAGGGACCAAGAGACCCCTAGCAGAGAGAGAGCTCCAGCTTCGTGTCATGATTC 220
QY 95 CCCAGTGCCTTCCTGCTGGGCTA 119
DB 221 ACCAGCTGTCCACCTCGGGACCA 245

RESULT 8

US-08-724-394A-20/c
Sequence 20, Application US/08724394A
Patent No. 5872237

GENERAL INFORMATION:
APPLICANT: Feder, John N.
APPLICANT: Krommal, Gregory S.
APPLICANT: Lauer, Peter M.
APPLICANT: Ruddy, David A.
APPLICANT: Thomas, Winston
APPLICANT: Tsuchihashi, Zenta
APPLICANT: Wolff, Roger K.
TITLE OF INVENTION: Megabase Transcript Map: No. 5872237el
TITLE OF INVENTION: Sequences and Antibodies Thereto
NUMBER OF SEQUENCES: 31
CORRESPONDENCE ADDRESS:
ADDRESS: TOWNSEND and TOWNSEND and CREW LLP
STREET: Two Embarcadero Center, 8th Floor
CITY: San Francisco
STATE: CA
COUNTRY: USA
ZIP: 94111-3834

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/724,394A
FILING DATE: 01-OCT-1996
CLASSIFICATION: 536

ATTORNEY/AGENT INFORMATION:
NAME: Fitts, Renee A.
REGISTRATION NUMBER: 35,136
REFERENCE/DOCKET NUMBER: 017957-000100
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-576-0200
TELEFAX: 415-576-0300
INFORMATION FOR SEQ ID NO: 20:
SEQUENCE CHARACTERISTICS:
LENGTH: 246240 base pairs
TYPE: nucleic acid
STRANDEDNESS: not relevant
TOPOLOGY: not relevant
MOLECULE TYPE: cDNA
FEATURE:
NAME/KEY: misc_feature
LOCATION: 1..246240

OTHER INFORMATION: /note= "HLA-H.CONTIG"
US-08-724-394A-20
Query Match 14.8%; Score 27; DB 2; Length 246240;
Best Local Similarity 62.7%; Pred. No. 54;
Matches 42; Conservative 0; Mismatches 25; Indels 0; Gaps 0;

Db 111423 TGCTTG 111417

Db 111483 CAGGAATGTAGTGGCAAGGAGAGAGAGGTAGTGTAATTCTCCTGCATCACCCCCC 111424

APPLICANT: Adler, Beverly
APPLICANT: Eragon Newell

;; TITLE OF INVENTION: N-Acetylglucosaminyltransferase V
;; TITLE OF INVENTION: Protein and Gene
;; NUMBER OF SEQUENCES: 19
;; CORRESPONDENCE ADDRESS:
;; ADDRESSEE: Greenlee and Winner, P.C.
;; STREET: 5370 Manhattan Circle, Suite 201
;; CITY: Boulder
;; STATE: Colorado
;; COUNTRY: USA
;; ZIP: 80303
;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: Floppy disk
;; COMPUTER: IBM PC compatible
;; OPERATING SYSTEM: PC-DOS/MS-DOS
;; SOFTWARE: Patentin Release #1.0, Version #1.25
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US 07/905,795
;; FILING DATE: 19930210
;; CLASSIFICATION: 435
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US 07/905,795
;; FILING DATE: 29-JUN-1992
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Ferber, Donna M.
;; REGISTRATION NUMBER: 33,878
;; REFERENCE/DOCKET NUMBER: 34-92A
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: (303)499-8080
;; TELEFAX: (303)499-8089
;; TELEX: 823189
;; INFORMATION FOR SEQ ID NO: 15:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 2624 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: double
;; TOPOLOGY: linear
;; MOLECULE TYPE: cDNA to mRNA
;; HYPOTHETICAL: NO
;; ANTI-SENSE: NO
;; FEATURE:
;; NAME/KEY: CDS
;; LOCATION: 299..2521
;; US-08-016-863-15

Query Match 14.5%; Score 26.6; DB 1; Length 2624;
Best Local Similarity 48.4%; Pred. No. 9.5;
Matches 74; Conservative 0; Mismatches 79; Indels 0; Gaps 0;
Db 9 CCCCCATTCTGAGCCCAAGAGGCTCATCCCTAAGGATGTCCAGAGATCCCAAGTGCAGAA 68
2489 CCTTGATGAAGTCCCGCAGGCGAGATCCGCTGGTGTGGGGTGGCCCGGCACAAC 2430
QY 69 GGAGATGTGGTGGAGGCTATTATTTCCCCAGTGCCTTCCCTGCTGGGCTATGGATGAAC 128
Db 2429 TGAAGACGAGGATCCCTTGGACACACAGTGTCTTCTTGGGTAGAGAGGGGCA 2370
QY 129 AGTGGTGAATTCATCTAGGAAGAGCTATGCG 161
Db 2369 CCAGGATGCTCTTGTATAGTTCTGAGCTTTGCG 2337

RESULT 12
US-08-276-968A-15/c
; Sequence 15, Application US/08276968A
; Patent No. 6015701
; GENERAL INFORMATION:
; APPLICANT: pierce, James M.
; APPLICANT: Shoreliah, Mohamed G.
; APPLICANT: Adler, Beverly L.
; APPLICANT: Fregien, Nevils L.
; TITLE OF INVENTION: N-Acetylglucosaminyltransferase V
; TITLE OF INVENTION: Proteins and Sequences
; NUMBER OF SEQUENCES: 34

;; CORRESPONDENCE ADDRESS:
;; ADDRESSEE: Greenlee and Winner, P.C.
;; STREET: 5370 Manhattan Circle, Suite 201
;; CITY: Boulder
;; STATE: Colorado
;; COUNTRY: USA
;; ZIP: 80303
;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: Floppy disk
;; COMPUTER: IBM PC compatible
;; OPERATING SYSTEM: PC-DOS/MS-DOS
;; SOFTWARE: Patentin Release #1.0, Version #1.30
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/08/276,968A
;; FILING DATE: 19-JUL-1994
;; CLASSIFICATION: 435
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US 07/905,795
;; FILING DATE: 29-JUN-1992
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US 08/016,863
;; FILING DATE: 10-FEB-1993
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Ferber, Donna M.
;; REGISTRATION NUMBER: 33,878
;; REFERENCE/DOCKET NUMBER: 34-92D
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: (303)-499-8080
;; TELEFAX: (303)-499-8089
;; TELEX: 49617824
;; INFORMATION FOR SEQ ID NO: 15:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 2624 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: double
;; TOPOLOGY: linear
;; MOLECULE TYPE: cDNA to mRNA
;; HYPOTHETICAL: NO
;; ANTI-SENSE: NO
;; FEATURE:
;; NAME/KEY: CDS
;; LOCATION: 299..2521
;; US-08-276-968A-15

Query Match 14.5%; Score 26.6; DB 3; Length 2624;
Best Local Similarity 48.4%; Pred. No. 9.5;
Matches 74; Conservative 0; Mismatches 79; Indels 0; Gaps 0;
QY 9 CCCCCATTCTGAGCCCAAGAGGCTCATCCCTAAGGATGTCCAGAGATCCCAAGTGCAGAA 68
Db 2489 CCTTGATGAAGTCCCGCAGGCGAGATCCGCTGGTGTGGGGTGGCCCGGCACAAC 2430
QY 69 GGAGATGTGGTGGAGGCTATTATTTCCCCAGTGCCTTCCCTGCTGGGCTATGGATGAAC 128
Db 2429 TGAAGACGAGGATCCCTTGGACACACAGTGTCTTCTTGGGTAGAGAGGGGCA 2370
QY 129 AGTGGTGAATTCATCTAGGAAGAGCTATGCG 161
Db 2369 CCAGGATGCTCTTGTATAGTTCTGAGCTTTGCG 2337

RESULT 13
US-09-434-408-1
; Sequence 1, Application US/09434408
; Patent No. 6440697
; GENERAL INFORMATION:
; APPLICANT: Venezia, Domenick
; APPLICANT: Grossmann, Angelika
; TITLE OF INVENTION: RING FINGER PROTEIN ZAFOP3
; FILE REFERENCE: 98-41
; CURRENT APPLICATION NUMBER: US/09/434,408
; CURRENT FILING DATE: 1999-11-04
; EARLIER APPLICATION NUMBER: US 60/108,258

EARLIER FILING DATE: 1998-11-12
NUMBER OF SEQ ID NOS: 23
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 1
LENGTH: 3138
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: CDS
LOCATION: (367)...(2535)
US-09-434-408-1

Query Match 14.2%; Score 26; DB 4; Length 3138;
Best Local Similarity 52.8%; Pred. No. 17;
Matches 56; Conservative 0; Mismatches 50; Indels 0; Gaps 0;
QY 8 TCCCCATTCTGAGCCCAAGAGCTCATCCCTAAGGATGTCAGAGATCCCAAGTGCAGA 67
DB 473 TCCTAATGTGAGCTCTCAGAGATTCATTGGAGCTTTGCAACATGCAAGTTCTGC 532
DB 68 AGGAGATGTGTGAGCTATTATTCCCTCCAGTGCCTTCCCTGCT 113
DB 533 AGAAGAGGTGCTGATGCTCCACAGAAATCACTCACTTCCCTGCT 578

RESULT 14
US-09-209-668-18
Sequence 18, Application US/09209668A
Patent No. 6114517
GENERAL INFORMATION:
APPLICANT: Monia, Brett P.
APPLICANT: Xu, Xiaoxing S.
TITLE OF INVENTION: METHODS OF MODULATING TUMOR NECROSIS FACTOR
TITLE OF INVENTION: alpha-INDUCED EXPRESSION OF CELL ADHESION MOLECULES
FILE REFERENCE: ISPH-0336
CURRENT APPLICATION NUMBER: US/09/209,668A
CURRENT FILING DATE: 1998-12-10
NUMBER OF SEQ ID NOS: 25
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 18
LENGTH: 3834
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: CDS
LOCATION: (117)...(1949)
PUBLICATION INFORMATION:
DATABASE ACCESSION NUMBER: M24736/Genbank
DATABASE ENTRY DATE: 1994-11-07
US-09-209-668-18

Query Match 14.2%; Score 26; DB 3; Length 3834;
Best Local Similarity 49.3%; Pred. No. 18;
Matches 68; Conservative 0; Mismatches 70; Indels 0; Gaps 0;
QY 21 GCCCAAGAGGCTCATCCCTAAGGATGTCAGAGATCCCAAGTGCAGAAGAGATGTGGT 80
DB 2072 GCTGAACCCGCAACACCCATCACCATTCAATAGATCAAAAGTCCAGCAGGACGGC 2131
QY 81 GAGGCTATTATTCCCTCCAGTGCCTTCCCTGCTGGGCTATGGATGACAGTGGCTGACTT 140
DB 2132 CTCAACTGAAAGAGACTCAGTGTTCCTTCTTCTACTCTCAGGATCAAGAAAGTGTGGCT 2191
QY 141 CATCTAGGAAAGAGCTAT 158
DB 2192 AATGAAGGGAAGGATAT 2209

RESULT 15
US-08-365-470-1
Sequence 1, Application US/08365470
Patent No. 5632991
GENERAL INFORMATION:

APPLICANT: Gimbrone, Jr., Michael A.
TITLE OF INVENTION: Antibodies Specific For E-selectin And The Uses
NUMBER OF SEQUENCES: 3
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX
STREET: 1100 New York Ave., NW
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/365,470
FILING DATE: herewith
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/102,510
FILING DATE: 05-AUG-1993
PRIOR APPLICATION DATA: US 07/850,802
FILING DATE: 13-MAR-1992
ATTORNEY/AGENT INFORMATION:
NAME: Markowicz, Karen R.
REGISTRATION NUMBER: 36,351
REFERENCE/DOCKET NUMBER: 0627.1350003
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 3854 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-365-470-1

Query Match 14.2%; Score 26; DB 1; Length 3854;
Best Local Similarity 49.3%; Pred. No. 18;
Matches 68; Conservative 0; Mismatches 70; Indels 0; Gaps 0;
QY 21 GCCCAAGAGGCTCATCCCTAAGGATGTCAGAGATCCCAAGTGCAGAAGAGATGTGGT 80
DB 2072 GCTGAACCCGCAACACCCATCACCATTCAATAGATCAAAAGTCCAGCAGGACGGC 2131
QY 81 GAGGCTATTATTCCCTCCAGTGCCTTCCCTGCTGGGCTATGGATGACAGTGGCTGACTT 140
DB 2132 CTCAACTGAAAGAGACTCAGTGTTCCTTCTTCTACTCTCAGGATCAAGAAAGTGTGGCT 2191
QY 141 CATCTAGGAAAGAGCTAT 158
DB 2192 AATGAAGGGAAGGATAT 2209

Search completed: May 25, 2003, 04:21:50
Job time : 354 secs

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OM nucleic - nucleic search, using sw model

Run on: May 25, 2003, 03:14:07 ; Search time 125 Seconds
(without alignments)
1933.157 Million cell updates/sec

Title: US-09-660-568-49
Perfect score: 183
Sequence: 1 cacacactcccccattctga.....ctgtctctggagctcacc 183

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 828747 seqs, 560231138 residues
Total number of hits satisfying chosen parameters: 1657494

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

- Database : Published Applications NA.*
- 1: /cgn2_5/ptodata/2/pubpna/US07_PUBCOMB.seq.*
 - 2: /cgn2_5/ptodata/2/pubpna/PCT_NEW_PUB.seq.*
 - 3: /cgn2_5/ptodata/2/pubpna/US06_NEW_PUB.seq.*
 - 4: /cgn2_5/ptodata/2/pubpna/US06_PUBCOMB.seq.*
 - 5: /cgn2_5/ptodata/2/pubpna/US07_NEW_PUB.seq.*
 - 6: /cgn2_5/ptodata/2/pubpna/PCTUS_PUBCOMB.seq.*
 - 7: /cgn2_5/ptodata/2/pubpna/US08_NEW_PUB.seq.*
 - 8: /cgn2_5/ptodata/2/pubpna/US08_PUBCOMB.seq.*
 - 9: /cgn2_5/ptodata/2/pubpna/US09_NEW_PUB.seq.*
 - 10: /cgn2_5/ptodata/2/pubpna/US09_PUBCOMB.seq.*
 - 11: /cgn2_5/ptodata/2/pubpna/US10_NEW_PUB.seq.*
 - 12: /cgn2_5/ptodata/2/pubpna/US10_PUBCOMB.seq.*
 - 13: /cgn2_5/ptodata/2/pubpna/US60_NEW_PUB.seq.*
 - 14: /cgn2_5/ptodata/2/pubpna/US60_PUBCOMB.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	35.2	19.2	164	10	US-09-864-761-21666
2	30.6	16.7	5216	9	US-10-125-540-607
3	30.6	16.7	5216	9	US-10-074-095-695
4	30.6	16.7	5216	10	US-09-764-870-607
5	30.6	16.7	5216	10	US-09-764-860-695
6	28.8	15.7	8894	9	US-10-092-154-1606
7	28.8	15.7	8894	10	US-09-764-847-1606
8	28.4	15.5	2001	9	US-09-829-155C-8
9	28.4	15.5	2513	9	US-09-829-155C-10
10	28.2	15.4	358	9	US-10-060-036-1937
11	28.2	15.4	1243	12	US-10-044-090-96
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13	28.2	15.4	2084	9	US-10-023-282-234
14	28.2	15.4	2329	12	US-10-044-090-416
15	27.8	15.2	374	9	US-09-918-995-6295
16	27.8	15.2	20907	9	US-09-764-891-9966
17	27.8	15.2	50000	9	US-10-152-724A-20
18	27.6	15.1	312	10	US-09-783-590-11431
19	27.6	15.1	945	9	US-10-023-601-43

20	27.6	15.1	1295	10	US-09-881-852-17	Sequence 17, Appl
21	27.6	15.1	26668	10	US-09-962-832-222	Sequence 222, App
22	27.6	15.1	39776	9	US-10-160-293-3	Sequence 3, Appl1
23	27.4	15.0	534	10	US-09-920-300A-1518	Sequence 1518, Ap
24	27.4	15.0	534	12	US-10-033-528-1518	Sequence 1518, Ap
25	27.4	15.0	904	9	US-10-082-830-120	Sequence 120, App
26	27.4	15.0	1309	9	US-10-082-830-121	Sequence 121, App
27	27.2	14.9	8048	9	US-09-764-872-917	Sequence 917, App
28	27.2	14.9	16825	9	US-10-092-154-1495	Sequence 1495, Ap
29	27.2	14.9	16825	10	US-09-764-847-1495	Sequence 1495, Ap
30	27	14.8	294	10	US-09-294-093B-1409	Sequence 1409, Ap
31	27	14.8	371	9	US-09-918-995-37556	Sequence 37556, A
32	27	14.8	422	10	US-09-920-300A-583	Sequence 583, App
33	27	14.8	422	12	US-10-033-528-583	Sequence 583, App
34	27	14.8	501	10	US-09-918-995-580-482	Sequence 482, App
35	27	14.8	502	9	US-09-918-995-32131	Sequence 32131, A
36	27	14.8	511	9	US-09-918-995-32726	Sequence 32726, A
37	27	14.8	56686	10	US-09-736-960-86	Sequence 86, Appl
38	26.8	14.6	1497	9	US-10-011-582-3	Sequence 3, Appl1
39	26.8	14.6	2175	9	US-10-011-582-1	Sequence 1, Appl1
40	26.8	14.6	2223	9	US-09-891-762-3	Sequence 3, Appl1
41	26.8	14.6	2286	9	US-10-011-582-9	Sequence 9, Appl1
42	26.8	14.6	2373	9	US-10-011-582-5	Sequence 5, Appl1
43	26.8	14.6	2612	9	US-10-011-582-8	Sequence 8, Appl1
44	26.8	14.6	4586	9	US-09-891-762-1	Sequence 1, Appl1
45	26.8	14.6	5746	9	US-09-764-891-5830	Sequence 5830, Ap

ALIGNMENTS

RESULT 1

US-09-864-761-21666

; Sequence 21666, Application US/09864761

; Patent No. US20020048763A1

; GENERAL INFORMATION:

; APPLICANT: Penn, Sharon G.

; APPLICANT: Rank, David R.

; APPLICANT: Hanzel, David K.

; APPLICANT: Chen, Wensheng

; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FO

; FILE REFERENCE: Acomica-x-1

; FILE REFERENCE: GENE EXPRESSION ANALYSIS BY MICROARRAY

; CURRENT APPLICATION NUMBER: US/09/864,761

; CURRENT FILING DATE: 2001-05-23

; PRIOR APPLICATION NUMBER: US 60/180,312

; PRIOR FILING DATE: 2000-02-04

; PRIOR APPLICATION NUMBER: US 60/207,456

; PRIOR FILING DATE: 2000-05-26

; PRIOR APPLICATION NUMBER: US 09/632,366

; PRIOR FILING DATE: 2000-08-03

; PRIOR APPLICATION NUMBER: GB 24263.6

; PRIOR FILING DATE: 2000-10-04

; PRIOR APPLICATION NUMBER: US 60/236,359

; PRIOR FILING DATE: 2000-09-27

; PRIOR APPLICATION NUMBER: PCT/US01/00666

; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00667

; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00664

; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00669

; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00665

; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00668

; PRIOR FILING DATE: 2001-01-30

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; PRIOR FILING DATE: 2001-01-30

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; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00661

; PRIOR FILING DATE: 2001-01-30

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; PRIOR APPLICATION NUMBER: PCT/US01/00670
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 60/234,687
; PRIOR FILING DATE: 2000-09-21
; PRIOR APPLICATION NUMBER: US 09/608,408
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: US 09/774,203
; PRIOR FILING DATE: 2001-01-29
; NUMBER OF SEQ ID NOS: 49117
; SOFTWARE: Anomax Sequence Listing Engine vers. 1.1
; SEQ ID NO 21666
; LENGTH: 164
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: MAP TO AF000498.1
; OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 1
; OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 1.1
; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.1
; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 0.97
; OTHER INFORMATION: EXPRESSED IN BT474, SIGNAL = 1
; OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 1
; OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 1.2
; OTHER INFORMATION: NT HIT: AB026898.1, EVALUE 9.00e-88
; OTHER INFORMATION: SWISSPROT HIT: P79023, EVALUE 1.70e-00
; OTHER INFORMATION: EST_HUMAN HIT: BF346467.1, EVALUE 1.40e-01
; US-09-864-761-21666

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Best Local Similarity 71.9%; Pred. No. 0.006; 18; Indels 0; Gaps 0;
Matches 46; Conservative 0; Mismatches 18; Indels 0; Gaps 0;

QY 52 GAGATCCAAAGTCAGAGGAGAGATGCTGAGGCTATTATTCCTCCAGTGCCTTCCTG 111
||||| ||||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||
Db 100 GAGATCAGAGACAGAGAGAGATGAGTGAATATTTATTCCTTGGCTCTCTCTTG 159

QY 112 CTGG 115
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Db 160 TGGG 163

RESULT 2
US-10-125-540-607/C
; Sequence 607, Application US/10125540
; Publication No. US20030059875A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PT214C1
; CURRENT APPLICATION NUMBER: US/10/125,540
; CURRENT FILING DATE: 2002-04-19
; Prior Application removed - See File Wrapper or Palm
; NUMBER OF SEQ ID NOS: 646
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 607
; LENGTH: 5216
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-10-125-540-607

Query Match          16.7%; Score 30.6; DB 9; Length 5216;
Best Local Similarity 56.4%; Pred. No. 0.92;
Matches 57; Conservative 0; Mismatches 44; Indels 0; Gaps 0;

QY 23 CCCAAGAGGCTATCCCTTAAGATGTCCAGAGATCCAGATCCAGAGAGGAGATGTGTA 82
||||| ||||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||
Db 2539 CTCGGAGGCTAAGGCAGGAGAGATGTGTGAACCCGGAAGGAGGAGGCAACTTGCAGT 2480

QY 83 GGCATTATTTCCCTCCAGTGCCTTCCCTCTGGCTATGGA 123
| || || || || || || || || || || || || || || || || || || || ||
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; PRIOR APPLICATION NUMBER: 09/764,860
; PRIOR FILING DATE: 2001-01-17
; PRIOR APPLICATION NUMBER: 60/179,065
; PRIOR FILING DATE: 2000-01-31
; PRIOR APPLICATION NUMBER: 60/180,628
; PRIOR FILING DATE: 2000-02-04
; PRIOR APPLICATION NUMBER: 60/214,886
; PRIOR FILING DATE: 2000-06-28
; PRIOR APPLICATION NUMBER: 60/217,487
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; PRIOR FILING DATE: 2000-08-14
; PRIOR APPLICATION NUMBER: 60/220,963
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; PRIOR APPLICATION NUMBER: 60/225,270
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; PRIOR APPLICATION NUMBER: 60/236,368
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; PRIOR FILING DATE: 2000-09-29
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; PRIOR APPLICATION NUMBER: 60/251,868
; PRIOR FILING DATE: 2000-12-08
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; PRIOR APPLICATION NUMBER: 60/229,287
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; PRIOR FILING DATE: 2000-09-05
; PRIOR APPLICATION NUMBER: 60/236,367
; PRIOR FILING DATE: 2000-09-29
; PRIOR APPLICATION NUMBER: 60/237,039
; PRIOR FILING DATE: 2000-10-02
; PRIOR APPLICATION NUMBER: 60/237,038
; PRIOR FILING DATE: 2000-10-02
; PRIOR APPLICATION NUMBER: 60/236,370
; PRIOR FILING DATE: 2000-09-29
; PRIOR APPLICATION NUMBER: 60/236,802
; PRIOR FILING DATE: 2000-10-02
; PRIOR APPLICATION NUMBER: 60/237,037
; PRIOR FILING DATE: 2000-10-02
; PRIOR APPLICATION NUMBER: 60/237,040
; PRIOR FILING DATE: 2000-10-02
; PRIOR APPLICATION NUMBER: 60/240,960
; PRIOR FILING DATE: 2000-10-20
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; PRIOR APPLICATION NUMBER: 60/239,937
; PRIOR FILING DATE: 2000-10-13
; PRIOR APPLICATION NUMBER: 60/241,787
; PRIOR FILING DATE: 2000-10-20
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; PRIOR FILING DATE: 2000-11-08
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; PRIOR FILING DATE: 2000-11-08
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; PRIOR FILING DATE: 2000-11-17
; PRIOR APPLICATION NUMBER: 60/249,210
; PRIOR FILING DATE: 2000-11-17
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; PRIOR FILING DATE: 2000-08-22
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; PRIOR APPLICATION NUMBER: 60/227,182
; PRIOR FILING DATE: 2000-08-22
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; PRIOR FILING DATE: 2000-08-14
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; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: 60/230,438
; PRIOR FILING DATE: 2000-09-06
; PRIOR APPLICATION NUMBER: 60/215,135
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; PRIOR FILING DATE: 2000-09-14
; PRIOR APPLICATION NUMBER: 60/232,397
; PRIOR FILING DATE: 2000-09-14
; PRIOR APPLICATION NUMBER: 60/232,399
; PRIOR FILING DATE: 2000-09-14
; PRIOR APPLICATION NUMBER: 60/232,401
; PRIOR FILING DATE: 2000-09-14
; PRIOR APPLICATION NUMBER: 60/241,808
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/241,826
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/241,786
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/241,221
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/246,475
; PRIOR FILING DATE: 2000-11-08
; PRIOR APPLICATION NUMBER: 60/231,243
; PRIOR FILING DATE: 2000-09-08

Query Match 16.7% Score 30.6; DB 9; Length 5216;
Best Local Similarity 56.4% Pred. No. 0.92;
Matches 57; Conservative 0; Mismatches 44; Indels 0; Gaps 0;

QY 23 CCCAAGAGGCTCATCCCTTAAGATGTCAGAGATCCAAAGTCAGAGAGAAATGTGTGA 82
DB 2678 CTCGGAGGCTAAGCAGGAGATGGTGTGAACCCGGAAGCAGAGGAGAACTTGCAGT 2737
QY 83 GGCTATTATTCCCTCCAGTGCCTCCCTGCTGGGCTATGGA 123
DB 2738 GAGCTTAGATTGCGCCACTGCACCTCCAGGCTGGGCAACAGA 2778

RESULT 4
US-09-764-870-607/c
; Sequence 607, Application US/09764870

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; Patent No. US20020042386A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PT214
; CURRENT APPLICATION NUMBER: US/09/764,870
; CURRENT FILING DATE: 2001-01-17
; Prior application data removed - consult PALM or file wrapper
; NUMBER OF SEQ ID NOS: 646
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 607
; LENGTH: 5216
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-764-870-607

Query Match
Best Local Similarity 16.7%; Score 30.6; DB 10; Length 5216;
Matches 57; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

23 CCCAAGAGGCTCATCCCTAAGGATGCCAGATCCAGATCCAGTCCAGAGGAGAATGTGGTGA 82
Db 2539 CTCGGGAGGCTAAGGCAGGAGAGATGTGTGAACCCGGAAGGCAGAGGCAAACTTGCACT 2480
QY 83 GGCATTATTTCCCTCCAGTGCCTTCCTGCTGGCTATGGA 123
Db 2479 GAGCTTAGATTGGCAGCTGCACCTCCAGGCTGGGCAACAGA 2439

RESULT 5
US-09-764-860-695
; Sequence 695, Application US/09764860
; Patent No. US20020094953A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PC008
; CURRENT APPLICATION NUMBER: US/09/764,860
; CURRENT FILING DATE: 2001-01-17
; Prior application data removed - consult PALM or file wrapper
; NUMBER OF SEQ ID NOS: 1198
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 695
; LENGTH: 5216
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-764-860-695

Query Match
Best Local Similarity 16.7%; Score 30.6; DB 10; Length 5216;
Matches 57; Conservative 0; Mismatches 44; Indels 0; Gaps 0;

QY 23 CCCAAGAGGCTCATCCCTAAGGATGCCAGATCCAGATCCAGTCCAGAGGAGAATGTGGTGA 82
Db 2678 CTCGGGAGGCTAAGGCAGGAGAGATGTGTGAACCCGGAAGGCAGAGGCAAACTTGCACT 2737
QY 83 GGCATTATTTCCCTCCAGTGCCTTCCTGCTGGCTATGGA 123
Db 2738 GAGCTTAGATTGGCAGCTGCACCTCCAGGCTGGGCAACAGA 2778

RESULT 6
US-10-092-154-1606
; Sequence 1606, Application US/10092154
; Publication No. US20030054375A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PC009C1
; CURRENT APPLICATION NUMBER: US/10/092,154
; CURRENT FILING DATE: 2002-03-07
; NUMBER OF SEQ ID NOS: 2003
; Prior Application removed - See File Wrapper or Palm

; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1606
; LENGTH: 8894
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-092-154-1606

Query Match
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Matches 51; Conservative 0; Mismatches 37; Indels 0; Gaps 0;

QY 56 GAAGGAGAATGTGGTGGAGCTATTATTTCCCCCGAGTGCCTCCCTGCTGGGTATGGATG 125
Db 3269 GAGGCAGAGGTTGTCAGTGAAGTATCGACCTGCACCTAGCCTGGGTGACAGAGC 3328
QY 126 AACAGTGGCTGACTTCATCTAGGAAAGA 153
Db 3329 GACAGAGCAGACTTCATCTAAGAAGA 3356

RESULT 7
US-09-764-847-1606
; Sequence 1606, Application US/09764847
; Patent No. US20020132767A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PC009
; CURRENT APPLICATION NUMBER: US/09/764,847
; CURRENT FILING DATE: 2001-01-17
; Prior application data removed - consult PALM or file wrapper
; NUMBER OF SEQ ID NOS: 2003
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1606
; LENGTH: 8894
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-764-847-1606

Query Match
Best Local Similarity 15.7%; Score 28.8; DB 10; Length 8894;
Matches 51; Conservative 0; Mismatches 37; Indels 0; Gaps 0;

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Db 3269 GAGGCAGAGGTTGTCAGTGAAGTATCGACCTGCACCTAGCCTGGGTGACAGAGC 3328
QY 126 AACAGTGGCTGACTTCATCTAGGAAAGA 153
Db 3329 GACAGAGCAGACTTCATCTAAGAAGA 3356

RESULT 8
US-09-829-155C-8/c
; Sequence 8, Application US/09829155C
; Patent No. US20020155561A1
; GENERAL INFORMATION:
; APPLICANT: Thayer, Edward C.
; TITLE OF INVENTION: Mammalian Disulfide Core Protein-4
; FILE REFERENCE: 00-29
; CURRENT APPLICATION NUMBER: US/09/829,155C
; CURRENT FILING DATE: 2002-04-03
; PRIOR APPLICATION NUMBER: 60/196,230
; PRIOR FILING DATE: 2000-04-10
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 8
; LENGTH: 2001
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-829-155C-8

Query Match
Best Local Similarity 15.5%; Score 28.4; DB 9; Length 2001;
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Db 245 TAGCTCCTCACTGCCTCTACTGGCCA 220

US-09-829-155C-10

Db 1221 TAGCTCCTCACTGCCCTCTACTGGCCA 1196

Query Match . 15.4%; Score 28.2; DB 9; Length 358;
Best Local Similarity 61.6%; Pred. No. 2.7;
Matches 45; Conservative 0; Mismatches 28; Indels

Query Match	15.4%	Score	28.2;	DB	9;	Length	358;
Best Local Similarity	61.6%	Pred. No.	2.7;				
Matches	45;	Conservative	0;	Mismatches	28;	Indels	0;
						Gaps	0;

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GenCore version 5.1.4_p5_4578
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OM nucleic - nucleic search, using sw model

Run on: May 25, 2003, 03:08:42 ; Search time 1933 Seconds
(without alignments)
2755.206 Million cell updates/sec

Title: US-09-660-568-49

Perfect score: 183

Sequence: 1 cacacactccccattctga.....ctgtctcttgagactcacca 183

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0

Maximum DB seq length: 20000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl:*

1: gb_ba:*

2: gb_htg:*

3: gb_in:*

4: gb_ov:*

5: gb_ov:*

6: gb_pat:*

7: gb_ph:*

8: gb_pl:*

9: gb_pr:*

10: gb_ro:*

11: gb_sts:*

12: gb_sy:*

13: gb_un:*

14: gb_vl:*

15: em_ba:*

16: em_fun:*

17: em_hum:*

18: em_in:*

19: em_mu:*

20: em_or:*

21: em_ov:*

22: em_pat:*

23: em_ph:*

24: em_pl:*

25: em_ro:*

26: em_sts:*

27: em_un:*

28: em_vl:*

29: em_vl:*

30: em_htg_hum:*

31: em_htg_inv:*

32: em_htg_other:*

33: em_htg_mus:*

34: em_htg_pln:*

35: em_htg_rod:*

36: em_htg_man:*

37: em_htg_vrt:*

38: em_sy:*

39: em_htgo_hum:*

40: em_htgo_mus:*

41: em_htgo_other:*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB	ID	Description
1	183	100.0	183	6	ARI30466	ARI30466 Sequence
C 2	152.2	83.2	146327	9	ALI162253	ALI162253 Human DNA
C 3	44.4	24.3	189509	9	AC099564	AC099564 Homo sapi
C 4	43.6	23.8	91639	9	ALI162399	ALI162399 Human DNA
C 5	43.4	23.7	167115	9	AC090093	AC090093 Homo sapi
C 6	43.4	23.7	173087	2	AC107318	AC107318 Sus scrofa
C 7	43.4	23.7	184787	2	AC087354	AC087354 Homo sapi
C 8	43.4	23.7	185420	2	AC021973	AC021973 Homo sapi
C 9	42.4	23.2	205816	9	AC013751	AC013751 Homo sapi
C 10	41.8	22.8	174380	2	AC016075	AC016075 Homo sapi
C 11	41.8	22.8	175559	9	AC112498	AC112498 Homo sapi
C 12	41.8	22.8	177220	9	HS179D22	HS179D22 Homo sapi
C 13	41.6	22.7	58693	2	AC107875	AC107875 Homo sapi
C 14	41.6	22.7	111461	9	AP005368	AP005368 Homo sapi
C 15	41.6	22.7	149126	9	AC105177	AC105177 Homo sapi
C 16	41.6	22.7	162987	2	AP005359	AP005359 Homo sapi
C 17	41.2	22.5	162126	9	AL354711	AL354711 Human DNA
C 18	41	22.4	60059	9	AL354657	AL354657 Human DNA
C 19	41	22.4	150409	9	AP003496	AP003496 Homo sapi
C 20	41	22.4	152142	9	AP003494	AP003494 Homo sapi
C 21	41	22.4	191896	9	AC009481	AC009481 Homo sapi
C 22	40.8	22.3	134339	9	AC114501	AC114501 Homo sapi
C 23	40.8	22.3	158513	2	AC011054	AC011054 Homo sapi
C 24	40.8	22.3	163395	2	AC103726	AC103726 Homo sapi
C 25	40.6	22.2	57000	9	AC078862	AC078862 Homo sapi
C 26	40.2	22.0	132470	9	HS203P18	HS203P18 Human DNA
C 27	40	21.9	150533	9	AL356215	AL356215 Human DNA
C 28	40	21.9	154067	2	AC087656	AC087656 Homo sapi
C 29	40	21.9	162851	2	AC009637	AC009637 Homo sapi
C 30	40	21.9	176425	2	AC090418	AC090418 Homo sapi
C 31	39.8	21.7	155147	9	HS992D9	HS992D9 Human DNA
C 32	39.8	21.7	162249	2	AC020682	AC020682 Homo sapi
C 33	39.8	21.7	166973	9	AL451049	AL451049 Human DNA
C 34	39.6	21.6	107637	2	AC094679	AC094679 Rattus no
C 35	39.6	21.6	147301	2	AP001928	AP001928 Homo sapi
C 36	39.6	21.6	167376	9	AP001994	AP001994 Homo sapi
C 37	39.6	21.6	180402	2	AC016849	AC016849 Homo sapi
C 38	39.6	21.6	221091	9	AP003390	AP003390 Homo sapi
C 39	39	21.3	64491	9	AL365188	AL365188 Human DNA
C 40	39	21.3	172900	9	AC107027	AC107027 Homo sapi
C 41	38.4	21.0	51626	2	AC068840	AC068840 Homo sapi
C 42	38.4	21.0	103120	2	AC103223	AC103223 Rattus no
C 43	38.4	21.0	112622	2	AC010866	AC010866 Homo sapi
C 44	38.4	21.0	114800	9	AC123567	AC123567 Homo sapi
C 45	38.4	21.0	119234	2	AC105231	AC105231 Homo sapi

ALIGNMENTS

RESULT 1
ARI30466 ARI30466 Sequence 49 from patent US 6190857. 183 bp DNA linear PAT 16-MAY-2001
LOCUS ARI30466
DEFINITION ARI30466
ACCESSION ARI30466
VERSION ARI30466.1 GI:14118791
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 183)
AUTHORS Ralph D., An G., O'Hara, S. Mark, and Veltri, R.
TITLE Diagnosis of disease state using mRNA profiles in peripheral leukocytes
JOURNAL Patent: US 6190857-A 49 20-FEB-2001;

FEATURES source Location/Qualifiers
1. 183
/organism="unknown"
BASE COUNT 43 a 52 c 45 g 43 t
ORIGIN

Query Match 100.0%; Score 183; DB 6; Length 183;
Best Local Similarity 100.0%; Pred. No. 3e-49;
Matches 183; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CACACATCCCCCATCTGAGCCCCAAGAGGCTCATCCCTAAGATGTCAGAGATCCAA 60
|||||
DB 1 CACACATCCCCCATCTGAGCCCCAAGAGGCTCATCCCTAAGATGTCAGAGATCCAA 60
|||||

QY 61 GTGCAGAGGAGAAATGTGTGAGGCTATTATTATCCCCAGTCCTCCCTGCTGGGCTAT 120
|||||
DB 61 GTGCAGAGGAGAAATGTGTGAGGCTATTATTATCCCCAGTCCTCCCTGCTGGGCTAT 120
|||||

QY 121 GGATGAACAGTGGCTGACATTCATCTAGGAAGAGCTATGGCTTCCTCTCGGAGCTCA 180
|||||
DB 121 GGATGAACAGTGGCTGACATTCATCTAGGAAGAGCTATGGCTTCCTCTCGGAGCTCA 180
|||||

QY 181 CCA 183
DB 181 CCA 183

RESULT 2
AL162253/c
LOCUS Human DNA sequence from clone RP11-574F11 on chromosome 9, complete
DEFINITION
ACCESSION AL162253
VERSION AL162253.17 GI:13677203
KEYWORDS HTG
SOURCE Homo sapiens
ORGANISM
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
AUTHORS Clark, G.
TITLE Direct Submission
JOURNAL Submitted (20-MAR-2001) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
COMMENT requests: clonerequests@sanger.ac.uk
On Apr 19, 2001 this sequence version replaced gi:12539553.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
This sequence was finished as follows unless otherwise noted: all
regions were either double-stranded or sequenced with an alternate
chemistry or covered by high quality data (i.e., phred quality >=
30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by at least
one plasmid subclone or more than one M13 subclone; and the
assembly was confirmed by restriction digest. The following
abbreviations are used to associate primary accession numbers given
in the feature table with their source databases: Em: EMBL; Sw:
SWISSPROT; Tr: TREMBL; Wp: WORMPEP; Information on the WORMPEP
database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep/ This sequence
was generated from part of bacterial clone contigs of human
chromosome 9, constructed by the Sanger Centre Chromosome 9 Mapping
Group. Further information can be found at
http://www.sanger.ac.uk/HGP/Chr9
RP11-574F11 is from the library RPCI-11.2 constructed by the group
of Pieter de Jong. For further details see
http://www.chori.org/bacpac/home.htm
VECTOR: pBACE3.6
IMPORTANT: This sequence is not the entire insert of clone

RP11-574F11 It may be shorter because we sequence overlapping
sections only once, except for a 100 base overlap.
The true right end of clone RP11-574F11 is at 146327 in this
sequence. The true left end of clone RP11-635N21 is at 62948 in
this sequence. The true right end of clone RP11-12D24 is at 100 in
this sequence.

FEATURES source Location/Qualifiers
1. 146327
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="9"
/clone="RP11-574F11"
/clone_lib="RPCI-11.2"
84304. 84605
misc_feature
/note="Sequence from overlapping clone BA635N21
(AL354744). Assembly confirmed by restriction digest
data."
84304. 84306
misc_feature
/note="Single clone region. Assembly confirmed by
restriction digest data."
BASE COUNT 44360 a 29767 c 29483 g 42717 t
ORIGIN

Query Match 83.2%; Score 152.2; DB 9; Length 146327;
Best Local Similarity 92.5%; Pred. No. 4.1e-39;
Matches 160; Conservative 0; Mismatches 13; Indels 0; Gaps 0;

QY 2 ACACATCCCCCATCTGAGCCCCAAGAGGCTCATCCCTAAGATGTCAGAGATCCAAAG 61
|||||
DB 145681 ACTCTCCACTCTGCTTGCAGCCCCAAGAGGCTCATCCCTAAGATGTCAGAGATCCAAAG 145622
|||||

QY 62 TGCAGAAGGAGAAATGTGTGAGGCTATTATTATCCCCAGTCCTCCCTGCTGGGCTATG 121
|||||
DB 145621 TGCAGAAGGAGAAATGTGTGAGGCTATTATTATCCCCAGTCCTCCCTGCTGGGCTATG 145562
|||||

QY 122 GATGAACAGTGGCTGACATTCATCTAGGAAGAGCTATGGCTTCCTCTCCCTGG 174
|||||
DB 145561 GATGAACAGTGGCTGACATTCATCTAGGAAGAGCTATGGCTTCCTCTCCCTGGTAG 145509
|||||

RESULT 3
AC099564/c
LOCUS Homo sapiens chromosome 1 clone RP11-359A17, complete sequence.
DEFINITION
AC099564 AL391540
ACCESSION AC099564.2 GI:20163088
VERSION
KEYWORDS HTG
SOURCE Homo sapiens
ORGANISM
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
AUTHORS Kaul, R.K., Olson, M.V., Zhou, Y., James, R.A., Rouse, G., Wu, Z.,
Saenphimmachak, C., Phelps, K.A., Raymond, C. and Haugen, E.D.
TITLE Direct Submission
JOURNAL Unpublished
COMMENT 2 (bases 1 to 189509)
REFERENCE Kaul, R.K., Olson, M.V., Raymond, C. and Haugen, E.D.
AUTHORS
TITLE Direct Submission
JOURNAL Submitted (16-NOV-2001) Genome Center, University of Washington,
Box 352145, Seattle, WA 98195, USA
REFERENCE 3 (bases 1 to 189509)
AUTHORS Kaul, R.K., Olson, M.V., Zhou, Y., James, R.A., Rouse, G., Wu, Z.,
Saenphimmachak, C., Phelps, K.A., Raymond, C. and Haugen, E.D.
TITLE Direct Submission
JOURNAL Submitted (17-APR-2002) Genome Center, University of Washington,
Box 352145, Seattle, WA 98195, USA
COMMENT On Apr 17, 2002 this sequence version replaced gi:16946001.
----- Genome Center
Center: University of Washington Genome Center
Center Code: UWGC
Web site: http://www.genome.washington.edu
Contact: uwgchgs@u.washington.edu

Drafting Center: SC	4125	4139	9625	10059	851	824
----- Project Information	-----	-----	-----	-----	-----	-----
Center project name: chr-1	6418	6369	4576	4663	8925	8862
Center clone name: RP11-359A17 (sc0138)	-----	-----	-----	-----	-----	-----
----- Summary Statistics	-----	-----	-----	-----	-----	-----
Sequencing vector: plasmid; L08752; 91% of reads	155	<800	5240	5198	5167	5132
Sequencing vector: plasmid; 9% of reads	-----	-----	-----	-----	-----	-----
Chemistry: Dye-terminator ET; 40% of reads	3110	3153	519	<800	4215	4171
Chemistry: Dye-terminator Big Dye; 60% of reads	-----	-----	-----	-----	-----	-----
Assembly program: Phrap; version 0.990319	335	<800	42	<800	998	1068
Consensus quality: 189243 bases at least Q40	-----	-----	-----	-----	-----	-----
Consensus quality: 189424 bases at least Q30	12736	12524	5005	4897	869	824
Consensus quality: 189508 bases at least Q20	-----	-----	-----	-----	-----	-----
Insert size: 189509; sum-of-contigs	429	<800	335	<800	674	<800
Quality coverage: 6.9x in Q20 bases; sum-of-contigs	-----	-----	-----	-----	-----	-----
-----	724	<800	1083	1078	432	<800
Overlapping Sequences:	-----	-----	-----	-----	-----	-----
5': Mapping in progress	1210	1212	1903	1876	1765	1759
3': RP11-455110 AL357499, 57084-bp overlap	-----	-----	-----	-----	-----	-----
-----	688	<800	7581	7291	2997	2956
Sequence Quality Assessment:	-----	-----	-----	-----	-----	-----
This entry has been annotated with sequence quality	2858	2845	277	<800	779	<800
estimates computed by the Phrap assembly program.	-----	-----	-----	-----	-----	-----
All manually edited bases have been reduced to quality zero.	6500	6369	1941	1876	3980	3919
Quality levels above 40 are expected to have less than	-----	-----	-----	-----	-----	-----
1 error in 10,000 bp.	115	<800	4444	4415	6832	7194
Base-by-base quality values are not generally visible from the	-----	-----	-----	-----	-----	-----
Genbank flat file format but are available as part	851	808	889	881	1239	1233
of this entry's ASN.1 file.	1104	1103	5285	5437	12095	12002
-----	-----	-----	-----	-----	-----	-----
This sequence was finished as follows unless otherwise noted:	15	<800	972	988	1532	1538
all regions were either double-stranded or sequenced with an	-----	-----	-----	-----	-----	-----
alternate chemistry or covered by high quality data (i.e., Phred	202	<800	11	<800	2586	2580
quality >= 30); an attempt was made to resolve all sequencing	-----	-----	-----	-----	-----	-----
problems, such as compressions and repeats; all regions were	10384	10323	4790	4897	6439	6380
covered by at least one plasmid subclone or more than one M13	-----	-----	-----	-----	-----	-----
subclone; and the assembly was confirmed by restriction digest.	660	<800	1180	1138	1250	1233
-----	-----	-----	-----	-----	-----	-----
Sequence Validation:	2553	2617	27590	27791	3830	3919
This sequence has been validated by Multiple Complete Digest	-----	-----	-----	-----	-----	-----
fingerprinting. Comparison of the experimentally derived digest	12237	11780	6201	6357	612	<800
fragments with sequence-predicted fragments is given below.	-----	-----	-----	-----	-----	-----
The electronically-digested sequence consists of both insert and	712	<800	2762	2751	2547	2580
vector, in order to accurately represent the entire circular BAC.	-----	-----	-----	-----	-----	-----
Small fragments below a variable cutoff (approximately 400-800 bp)	2156	2175	351	<800	6491	6380
are not resolved in the fingerprint and hence do not appear	-----	-----	-----	-----	-----	-----
in the table. There are no significant remaining discrepancies	559	<800	3383	3355	553	<800
between the experimental and predicted values. Uniquely ordered	-----	-----	-----	-----	-----	-----
fragments are separated by dashed lines.	12040	11780	13408	12546	475	<800
-----	-----	-----	-----	-----	-----	-----
EcoRI	-----	-----	-----	-----	-----	-----
SeqDerMap FngPrnt SeqDerMap FngPrnt SeqDerMap FngPrnt	1752	1756	5980	6048	1993	1946
-----	-----	-----	-----	-----	-----	-----
BglII	6078	6013	1006	1078	3063	3123
-----	-----	-----	-----	-----	-----	-----
HindIII	1736	1756	9250	9321	5043	4966
-----	-----	-----	-----	-----	-----	-----
SeqDerMap FngPrnt SeqDerMap FngPrnt SeqDerMap FngPrnt	3432	3387	6101	6048	4148	4171
-----	-----	-----	-----	-----	-----	-----
8696	8707	2944	2929	529	<800	<800
-----	-----	-----	-----	-----	-----	-----
6	<800	2067	2050	6382	6380	6380
-----	-----	-----	-----	-----	-----	-----
6553	6369	6296	6357	512	<800	<800
-----	-----	-----	-----	-----	-----	-----
3847	3850	1756	1717	449	<800	<800
-----	-----	-----	-----	-----	-----	-----
1347	1329	1273	1295	3402	3320	3320
-----	-----	-----	-----	-----	-----	-----
1619	1607	199	<800	2513	2580	2580
-----	-----	-----	-----	-----	-----	-----
1509	1483	1163	1138	7289	7194	7194
-----	-----	-----	-----	-----	-----	-----
734	<800	5058	5198	1890	1946	1946
-----	-----	-----	-----	-----	-----	-----
96	<800	5880	5768	1165	1134	1134
-----	-----	-----	-----	-----	-----	-----
639	<800	1593	1555	2969	2956	2956

Roy, A., Santos, R., Schauer, S., Schupbach, R., Seaman, S., Severy, P., Sougnuez, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Testaye, S., Travers, M., Topham, K., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

TITLE
JOURNAL
REFERENCE
AUTHORS

Direct Submission
Submitted (14-FEB-2003) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
(bases 1 to 167115)

Birren, B., Linton, L., Nusbaum, C., Lander, E., Ali, A., Allen, N., Anderson, S., Barna, N., Bastien, V., Bloom, T., Boguslavsky, L., Boukhalter, B., Brown, A., Camarata, J., Campopiano, A., Chang, J., Chazaro, B., Choepel, Y., Colangelo, M., Collins, S., Collymore, A., Cook, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J. S., Dodge, S., Faro, S., Ferreira, P., FitzHugh, W., Gage, D., Galagan, J., Gardyna, S., Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-Pierre, N., Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., LaRocque, K., Lamazares, R., Landers, R., Lehoczy, J., Levine, R., Lindblad-Toh, K., Liu, G., MacLean, C., Macdonald, P., Major, J., Marquis, N., Matthews, C., McCarthy, M., McEwan, P., McKernan, K., Meldrim, J., Meneus, L., Mihova, T., Norman, C. H., O'Connor, T., O'Donnell, P., O'Neil, D., Oliver, J., Peterson, K., Phunkhang, P., Rieback, M., Riley, R., Rise, C., Rogov, P., Rosetti, M., Roy, A., Santos, R., Schauer, S., Schupbach, R., Seaman, S., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N., Strauss, N., Subramanian, A., Talamas, J., Testaye, S., Theodore, J., Topham, K., Travers, M., Travis, N., Trigilio, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

TITLE
JOURNAL
REFERENCE
AUTHORS

Direct Submission
Submitted (02-MAY-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
(bases 1 to 167115)

Birren, B., Linton, L., Nusbaum, C., Lander, E., Ali, A., Allen, N., Anderson, S., Barna, N., Bastien, V., Bloom, T., Boguslavsky, L., Boukhalter, B., Brown, A., Camarata, J., Campopiano, A., Chang, J., Chazaro, B., Choepel, Y., Colangelo, M., Collins, S., Collymore, A., Cook, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J. S., Dodge, S., Faro, S., Ferreira, P., FitzGerald, M., FitzHugh, W., Gage, D., Galagan, J., Gardyna, S., Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-Pierre, N., Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., LaRocque, K., Lamazares, R., Landers, R., Lehoczy, J., Levine, R., Lindblad-Toh, K., Liu, G., MacLean, C., Macdonald, P., Major, J., Marquis, N., Matthews, C., McCarthy, M., McEwan, P., McKernan, K., Meldrim, J., Meneus, L., Mihova, T., Mlenka, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., Norman, C. H., O'Connor, T., O'Donnell, P., O'Neil, D., Oliver, J., Peterson, K., Phunkhang, P., Rieback, M., Riley, R., Rise, C., Rogov, P., Roman, J., Rosetti, M., Roy, A., Santos, R., Schauer, S., Schupbach, R., Seaman, S., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N., Strauss, N., Subramanian, A., Talamas, J., Testaye, S., Theodore, J., Topham, K., Travers, M., Travis, N., Trigilio, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

TITLE
JOURNAL
REFERENCE
AUTHORS

Direct Submission
Submitted (31-MAY-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
(bases 1 to 167115)

Birren, B., Linton, L., Nusbaum, C., Lander, E., Ali, A., Allen, N., Anderson, S., Barna, N., Bastien, V., Bloom, T., Boguslavsky, L., Boukhalter, B., Brown, A., Camarata, J., Campopiano, A., Chang, J., Chazaro, B., Choepel, Y., Colangelo, M., Collins, S., Collymore, A., Cook, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J. S., Dodge, S., Faro, S., Ferreira, P., FitzGerald, M., FitzHugh, W., Gage, D., Galagan, J., Gardyna, S., Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-Pierre, N., Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., LaRocque, K., Lamazares, R., Landers, R., Lehoczy, J., Levine, R., Lindblad-Toh, K., Liu, G., MacLean, C., Macdonald, P., Major, J., Marquis, N.,

Matthews, C., McCarthy, M., McEwan, P., McKernan, K., Meldrim, J., Meneus, L., Mihova, T., Mlenka, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., Norman, C. H., O'Connor, T., O'Donnell, P., O'Neil, D., Oliver, J., Peterson, K., Phunkhang, P., Rieback, M., Riley, R., Rise, C., Pollara, V., Raymond, C., Retta, R., Rieback, M., Riley, R., Rise, C., Rogov, P., Roman, J., Rosetti, M., Roy, A., Santos, R., Schauer, S., Schupbach, R., Seaman, S., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N., Strauss, N., Subramanian, A., Talamas, J., Testaye, S., Theodore, J., Topham, K., Travers, M., Travis, N., Trigilio, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

Direct Submission

Submitted (01-JUN-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
On May 31, 2002 this sequence version replaced gi:20389333.

All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WtBR
Web site: <http://www-seq.wi.mit.edu>
Contact: sequence.submissions@genome.wi.mit.edu

----- Project Information
Center project name: L12753
Center clone name: 656_A_10

----- Location/Qualifiers
1. 167115
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="8"
/map="8"

----- Features
source
unsure
unsure
unsure
unsure
repeat_region
unsure
unsure
unsure
unsure
repeat_region
repeat_region
repeat_region
repeat_region
repeat_region
repeat_region
repeat_region
repeat_region

note="single clone coverage"
note="<30 qual SINGL region"
note="<30 qual SINGL region"
note="<30 qual SINGL region"
note="<30 qual SINGL region"
note="<30 qual SINGL region"
note="<30 qual SINGL region"
note="<30 qual SINGL region"
note="<30 qual SINGL region"
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note="<30 qual SINGL region"
note="<30 qual SINGL region"
note="<30 qual SINGL region"
note="<30 qual SINGL region"
note="<30 qual SINGL region"
note="<30 qual SINGL region"
note="<30 qual SINGL region"

clone="RP11-656A10"
clone.lib="RPC1-11 Human Male BAC"
374..393
471..475
508..512
572..576
complement(642..1043)
rpt_family="MER66A"
985..990
note="<30 qual SINGL region"
1042..1046
1062..1067
1075..1079
1102..1110
1121..1130
complement(1222..1485)
rpt_family="ERVL"
complement(1551..1852)
rpt_family="ERVL"
complement(2187..2378)
rpt_family="L3"
2530..2923
rpt_family="L2"
4193..4411
rpt_family="AluSx"
4844..4961
rpt_family="MIR"
complement(4950..5023)
rpt_family="MIR3"
complement(5475..5561)
rpt_family="LIPB3"

```

----- Project Information
Center project name: caz
Center clone name: 137G11
----- Summary Statistics
Sequencing method: plasmid; n/a; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 171731 bases at least Q40
Consensus quality: 172220 bases at least Q30
Consensus quality: 172357 bases at least Q20
Insert size: 143000; agarose-fp
Insert size: 172787; sum-of-contigs
Quality coverage: 11.24x in Q20 bases; agarose-fp
Quality coverage: 9.30x in Q20 bases; sum-of-contigs
-----
** NOTE: This is a 'working draft' sequence. It currently
** consists of 4 contigs. The true order of the pieces
** is not known and their order in this sequence record is
** arbitrary. Gaps between the contigs are represented as
** runs of N, but the exact sizes of the gaps are unknown.
** This record will be updated with the finished sequence
** as soon as it is available and the accession number will
** be preserved.
**
** 1 16132: contig of 16132 bp in length
** 16133 16232: gap of unknown length
** 16233 33937: contig of 17705 bp in length
** 33938 34037: gap of unknown length
** 34038 98779: contig of 64742 bp in length
** 98780 98879: gap of unknown length
** 98880 173087: contig of 74208 bp in length.
Location/Qualifiers
1..173087
/organism="Sus scrofa"
/db_xref="taxon:9823"
/clone="RP44-137G11"
/clone_lib="RP44"
1..16132
/note="assembly_fragment"
16233..33937
/note="assembly_fragment"
34038..98779
/note="assembly_fragment"
clone_end:SP6
vector_side:left"
98880..173087
/note="assembly_fragment"
clone_end:T7
vector_side:left"
52234 a 32792 c 33848 g 53909 t 304 others
              23.7%; Score 43.4; DB 2; Length 173087;
Similarity 65.3%; Pred.No.0.0029;
79; Conservative 0; Mismatches 41; Indels 1; Gaps 1;
          TCCAGAGATCAAGTCAGACGAGAAGTGTGGTGAGCTATTATTTCCCAGTCGCTT 106
          ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
          CCAGAGGTCCAGGTGCAGACGAGGGTGAGGTAGGTATTTATTTCTTGTTTC-T 15630
          CTGCTGGGCTATGGATCAACACTGGCTGACTTCATCTAGGAAGAGCATGCTTCTG 166
          ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
          CCTGCCAGGCTGTGGTTAGCAGTGGTTGTTCTTCTTACTCTCAATTTCTGGCTTCTC 15690
167
15691

```


Insert size: 185000; agarose-fp
 Insert size: 184820; sum-of-contigs
 Quality coverage: 6.44 in Q20 bases; agarose-fp
 Quality coverage: 5.98 in Q20 bases; sum-of-contigs

 * NOTE: This is a 'working draft' sequence. It currently
 * consists of 7 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

1 2842: contig of 2842 bp in length
 2843 2942: contig of unknown length
 2943 6917: contig of 3975 bp in length
 6918 7017: gap of unknown length
 7018 15680: contig of 8663 bp in length
 15681 15780: gap of unknown length
 15781 30574: contig of 14794 bp in length
 30575 30674: gap of unknown length
 30675 61322: contig of 30648 bp in length
 61323 61422: gap of unknown length
 61423 93101: contig of 31679 bp in length
 93102 93201: gap of unknown length
 93202 185420: contig of 92219 bp in length.

FEATURES

Source

1. 185420
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /chromosome="8"
 /clone="RP11-198H6"

misc_feature

1. 2842
 /note="assembly_name:Contig8
 clone_end:77
 vector_side:right"

misc_feature

2943..6917

misc_feature

/note="assembly_name:Contig9"

misc_feature

7018..15680

misc_feature

/note="assembly_name:Contig10"

misc_feature

15781..30574

misc_feature

/note="assembly_name:Contig11"

misc_feature

30675..61322

misc_feature

/note="assembly_name:Contig12"

misc_feature

61423..93101

misc_feature

/note="assembly_name:Contig13
 clone_end:SP6
 vector_side:left"

misc_feature

93202..185420

BASE COUNT

53244 a 39520 c 39207 g 52848 t 601 others

ORIGIN

Query Match 23.7%; Score 43.4; DB 2; Length 185420;
 Best Local Similarity 72.7%; Pred. No. 0.0029;
 Matches 56; Conservative 0; Mismatches 21; Indels 0; Gaps 0;

QY 65 AGAAGGAGAAATGTTGGTGGAGCTATTATTCCTCCAGTGCCTCCCTGGGTATGGAT 124

Db 167596 AGAAGCAGAGAGAGATCAGGCGCTTTGTTACCAAGTTCCTCCCTCTCTAAGGTT 167537

QY 125 GACAGTGGCTGACTTC 141

Db 167536 TCACAGTGGCTTTCTTC 167520

RESULT 9

AC013751

LOCUS

DEFINITION

AC013751

ACCESSION

VERSION

KEYWORDS

AC013751 AC013751 205816 bp DNA linear PRI 07-MAY-2001
 DEFINITION Homo sapiens chromosome, clone RP11-298P6, complete sequence.
 AC013751
 AC013751.6 GI:13958504
 HTG.

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

Homo sapiens.

Homo sapiens

Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

1 (bases 1 to 205816)

Birren,B., Linton,L., Nusbaum,C. and Lander,E.

Homo sapiens chromosome, clone RP11-298P6

Unpublished

2 (bases 1 to 205816)

Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,

Baldwin,J., Barna,N., Beckerly,R., Boguslavsky,L., Boukhgalter,B.,

Brown,A., Castelle,A., Colangelo,M., Collins,S., Collymore,A.,

Cooke,P., Dearellano,K., Dewar,K., Domino,M., Doneilan,L., Doyle,M.,

Ferreira,P., FitzHugh,W., Forrest,C., Funke,R., Gage,D.,

Galagan,J., Gardyna,S., Grant,G., Hagos,B., Hearford,A., Horton,L.,

Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J.,

Lehoczy,J., Liew,C., Locke,K., Macdonald,P., Marquis,N.,

McEwan,P., McGurk,A., McKernan,K., McLaughlin,J., Meldrim,J.,

Morrow,J., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,

Peterson,K., Pollara,V., Riley,R., Roy,A., Santos,R., Severy,P.,

Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,

Tesfaye,S., Tirrell,A., Vassiliev,H., Vo,A., Wheeler,J., Wu,X.,

Wyman,D., Ye,W.J., Zimmer,A. and Zody,M.

Direct Submission

Submitted (15-NOV-1999) Whitehead Institute/MIT Center for Genome

Research, 320 Charles Street, Cambridge, MA 02141, USA

3 (bases 1 to 205816)

Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,S.,

Barna,N., Bastien,V., Boguslavsky,L., Boukhgalter,B., Brown,A.,

Camarata,J., Campopiano,A., Chang,J., Choepel,Y., Colangelo,M.,

Collins,S., Collymore,A., Cooke,P., Dearellano,K., Dewar,K.,

Diaz,J.S., Dodge,S., Faro,S., Ferreira,P., FitzHugh,W., Gage,D.,

Galagan,J., Gardyna,S., Ginde,S., Goyette,M., Graham,L.,

Grand-Pierre,N., Hagos,B., Hearford,A., Horton,L., Hulme,W.,

Iliev,I., Johnson,R., Jones,C., Karatas,A., LaRoque,K.,

Lamazares,R., Landers,T., Lehoczy,J., Levine,R., Liu,G.,

Maclean,C., Macdonald,P., Marquis,N., Matthews,C., McCarthy,M.,

McEwan,P., McKernan,K., McPheeters,R., Meldrim,J., Meneus,L.,

Mihova,T., Mienga,V., Murphy,T., Naylor,J., Nguyen,C., Norbu,C.,

Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J.,

Peterson,K., Phunkhang,P., Pierre,N., Pollara,V., Raymond,C.,

Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P., Roman,J.,

Rosetti,M., Roy,A., Santos,R., Schauer,S., Schupback,R., Seaman,S.,

Severy,P., Sougnez,C., Spencer,B., Stange-Thomann,N.,

Stojanovic,N., Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S.,

Theodore,J., Travers,M., Travis,N., Trigilio,J., Vassiliev,H.,

Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.,

Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

Direct Submission

Submitted (06-MAY-2001) Whitehead Institute/MIT Center for Genome

Research, 320 Charles Street, Cambridge, MA 02141, USA

4 (bases 1 to 205816)

Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,S.,

Barna,N., Bastien,V., Boguslavsky,L., Boukhgalter,B., Brown,A.,

Camarata,J., Campopiano,A., Chang,J., Choepel,Y., Colangelo,M.,

Collins,S., Collymore,A., Cooke,P., Dearellano,K., Dewar,K.,

Diaz,J.S., Dodge,S., Faro,S., Ferreira,P., FitzHugh,W., Gage,D.,

Galagan,J., Gardyna,S., Ginde,S., Goyette,M., Graham,L.,

Grand-Pierre,N., Hagos,B., Hearford,A., Horton,L., Hulme,W.,

Iliev,I., Johnson,R., Jones,C., Karatas,A., LaRoque,K.,

Lamazares,R., Landers,T., Lehoczy,J., Levine,R., Liu,G.,

Maclean,C., Macdonald,P., Marquis,N., Matthews,C., McCarthy,M.,

McEwan,P., McKernan,K., McPheeters,R., Meldrim,J., Meneus,L.,

Mihova,T., Mienga,V., Murphy,T., Naylor,J., Nguyen,C., Norbu,C.,

Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J.,

Peterson,K., Phunkhang,P., Pierre,N., Pollara,V., Raymond,C.,

Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P., Roman,J.,

Rosetti,M., Roy,A., Santos,R., Schauer,S., Schupback,R., Seaman,S.,

Severy,P., Sougnez,C., Spencer,B., Stange-Thomann,N.,

Stojanovic,N., Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S.,

Theodore,J., Travers,M., Travis,N., Trigilio,J., Vassiliev,H.,

Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.,

Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

VERSION KEYWORDS SOURCE ORGANISM

AC016075.4 GI:7259724
HTG: HTGS_PHASE1; HTGS_DRAFT.

Homo sapiens

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Euthera; Primates; Catarrhini; Homnidae; Homo.

1 (bases 1 to 174380)

Birren,B., Linton,L., Nusbaum,C. and Lander,E.

Homo sapiens, clone RP11-699A7

Unpublished

2 (bases 1 to 174380)

Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,
Baldwin,J., Barna,N., Beckerly,R., Boguslavsky,L., Boukhgalter,B.,
Brown,A., Castelle,A., Collange,M., Collins,S., Collymore,A.,
Cooke,P., Dearellano,K., Dewar,K., Domino,M., Donegan,L., Doyle,M.,
Farreira,P., FitzHugh,W., Forrest,C., Funke,R., Gage,D.,
Galagan,J., Gardyna,S., Grant,G., Hagos,B., Heaford,A., Horton,L.,
Howland,J.C., Johnson,R., Jones,C., Kano,L., Karatas,A., Klein,J.,
Lehoczy,J., Lieu,C., Locke,K., Macdonald,P., Marquis,N.,
McEwan,P., McGurk,A., McKernan,K., McLaughlin,J., Meldrum,J.,
Morrow,J., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,
Peterson,K., Pollara,V., Riley,R., Roy,A., Santos,R., Severy,P.,
Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,
Tesfaye,S., Tirell,A., Vassiliev,H., Vo,A., Wheeler,J., Wu,X.,
Wyman,D., Ye,W.J., Zimmer,A. and Zody,M.

TITLE JOURNAL

Submitted (20-NOV-1999) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA

On Mar 17, 2000 this sequence version replaced gi:6970517.

All repeats were identified using RepeatMasker:

Smit, A.F.A. & Green, P. (1996-1997)

http://ftp.genome.washington.edu/RM/RepeatMasker.html

----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBR

Web site: http://www-seq.wi.mit.edu

Contact: sequence_submissions@genome.wi.mit.edu

----- Project Information

Center project name: L4345

Center clone name: 699_A7

----- Summary Statistics

Sequencing vector: M13; M77815; 100% of reads

Chemistry: Dye-terminator Big Dye; 100% of reads

Assembly program: Phrap; version 0.960731

Consensus quality: 166563 bases at least Q40

Consensus quality: 169997 bases at least Q30

Consensus quality: 171469 bases at least Q20

Insert size: 181000; agarose-fp

Insert size: 173280; sum-of-contigs

Quality coverage: 5.2 in Q20 bases; agarose-fp

Quality coverage: 5.4 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 12 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

* 1 377: contig of 377 bp in length
* 378 477: gap of 100 bp
* 478 1740: contig of 1263 bp in length
* 1741 1840: gap of 100 bp
* 1841 4031: contig of 2191 bp in length
* 4032 4131: gap of 100 bp
* 4132 9969: contig of 5838 bp in length
* 9970 10069: gap of 100 bp
* 10070 17346: contig of 7277 bp in length
* 17347 17446: gap of 100 bp
* 17447 29159: contig of 11713 bp in length
* 29160 29259: gap of 100 bp
* 29260 44992: contig of 15733 bp in length

FEATURES source

* 44993 45092: gap of 100 bp
* 45093 63565: contig of 18473 bp in length
* 63566 63665: gap of 100 bp
* 63666 82201: contig of 18536 bp in length
* 82202 82301: gap of 100 bp
* 82302 105742: contig of 23441 bp in length
* 105743 105842: gap of 100 bp
* 105843 139307: contig of 33465 bp in length
* 139308 139407: gap of 100 bp
* 139408 174380: contig of 34973 bp in length.

Location/Qualifiers

1..174380
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="RP11-699A7"

1..377
/note="assembly_fragment"

clone_end:17
vector_side:right

478..1740
/note="assembly_fragment"

1841..4031
/note="assembly_fragment"

4132..9969
/note="assembly_fragment"

10070..17346
/note="assembly_fragment"

17447..29159
/note="assembly_fragment"

29260..44992
/note="assembly_fragment"

45093..63565
/note="assembly_fragment"

63666..82201
/note="assembly_fragment"

clone_end:SP6
vector_side:left

82302..105742
/note="assembly_fragment"

105843..139307
/note="assembly_fragment"

139408..174380
/note="assembly_fragment"

BASE COUNT 53438 a 32639 c 33542 g 53658 t 1103 others

ORIGIN

Query Match 22.8%; Score 41.8; DB 2; Length 174380;
Best Local Similarity 63.4%; Pred. No. 0.0096;
Matches 64; Conservative 0; Mismatches 37; Indels 0; Gaps 0;
QY 52 GAGATCCAAAGTCAGAGAGAGATGTGGTGGAGGCTATTTATTTCCCGCAGTGCCTTCCCTG 111
DB 154021 GAGATTGGAGACAGAGAATAGCAAGATCATGATATTTGTTCCCTTAGCTACATCCATG 154080
QY 112 CTGGGCTATGGATGACACTGGCTGACTTCATCTAGGAAG 152
DB 154081 CAGGGCCACTGGTGGCAGTGTCTCTTCTCTAGTAAGG 154121

RESULT 11

AC112498/c

LOCUS

DEFINITION

Homo sapiens X BAC RP11-699A7 (Roswell Park Cancer Institute Human

BAC Library) complete sequence.

ACCESSION

AC112498

VERSION

AC112498.3 GI:21591809

KEYWORDS

HTG.

SOURCE

human.

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Euthera; Primates; Catarrhini; Homnidae; Homo.

REFERENCE

1 (bases 1 to 175559)

AUTHORS

Muzny, D.M., Adams, C., Adio-Oduola, B., Ali-Osman, F.R., Allen, C., Alsbrooks, S.L., Amarantunga, H.C., Are, J.R., Banks, T., Barbaria, J., Benton, J., Bimaga, K., Blankenburg, K., Bonnin, D., Bouck, J., Bowie, S., Brieva, M., Brown, E., Brown, M., Bryant, N.P., Buhay, C., Burkett, C., Burrell, K.L., Byrd, N.C., Carton, T.F., Carter, M., Cavazos, S.R., Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Z., Chowdhury, I., Christopoulos, C., Cleveland, C.D., Cox, C., Coyle, M.D., Dathorne, S.R., David, R., Davila, M.L., Davis, C., Davy-Carroll, L., Dederich, D.A., Delaney, K.R., Delgado, O., Denn-A-Li, Ding, Y., Dinh, H.H., Douthwaite, K.J., Draper, H., Dugan-Rocha, S., Durbin, K.J., Earnhart, C., Edgar, D., Edwards, C.C., Elhaj, C., Escotto, M., Falls, T., Ferraguto, D., Flagg, N., Ford, J., Foster, P., Frantz, P., Gabisi, A., Gao, J., Garcia, A., Garner, T., Garza, N., Gill, R., Gorrell, J.H., Guevara, W., Gunaratne, P., Hale, S., Hamilton, K., Harris, C., Harris, K., Hart, M., Havlak, P., Hawes, A., Hernandez, J., Hernandez, O., Hodgson, A., Hoques, M., Holloway, C., Hollins, B., Honsi, F., Howard, S., Huber, J., Hulyk, S., Hume, J., Jackson, L.E., Jacobson, B., Jia, Y., Johnson, R., Jolivet, S., Joudah, S., Karlsson, E., Kelly, S., Khan, U., King, L., Korvah, J., Kovar, C., Kratovic, J., Kureshi, A., Landry, N., Leal, B., Lewis, L.C., Lewis, L., Li, J., Li, Z., Lichtarge, O., Lieu, C., Liu, J., Liu, W., Loulsegh, H., Lozador, R.J., Lu, X., Lucier, A., Lucier, R., Luna, R., Ma, J., Maheshwari, M., Mapua, P., Martin, R., Martindale, A., Martinez, E., Massey, E., Mawhiney, E., McLeod, M.P., Meador, M., Mei, G., Metzker, M., Miner, G., Miner, Z., Mitchell, T., Mohabbat, K., Morgan, M., Morris, S., Moser, M., Neal, D., Newton, J., Newton, N., Nguyen, A., Nguyen, N., Nguyen, N., Nickerson, E., Nwokenkwo, S., Ogih, M., Okuwonu, G., Oragunye, N., Oviedo, R., Pace, A., Payton, B., Peery, J., Perez, L., Peters, L., Pickens, R., Primus, E., Pu, L.L., Quiles, M., Ren, Y., Rives, M., Rojas, A., Rojibokan, I., Rolfe, M., Ruiz, S., Savery, G., Scherer, S., Scott, G., Shen, H., Shooshtari, N., Sisson, I., Sodergren, E., Sonaite, T., Sparks, A., Stanley, H., Stone, H., Sutton, A., Svatek, A., Tabor, P., Tamerisa, A., Tamerisa, K., Tang, H., Tansey, J., Taylor, C., Taylor, T., Telford, B., Thomas, N., Thomas, S., Usmani, K., Vasquez, L., Vera, V., Villalón, D., Vinson, R., Wang, Q., Wang, S., Ward-Moore, S., Warren, R., Washington, C., Watlington, S., Williams, G., Williamson, A., Wleczyk, R., Wooden, S., Worley, K., Wu, C., Wu, F., Wu, F., Zhou, J., Zorrilla, S., Nelson, D., Weinstock, G. and Gibbs, R.

Direct Submission
Unpublished
2 (bases 1 to 175559)
Worley, K.C.
Direct Submission
Submitted (21-FEB-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 175559)
Worley, K.C.
Direct Submission
Submitted (09-MAY-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
4 (bases 1 to 175559)
Worley, K.C.
Direct Submission
Submitted (26-JUN-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
5 (bases 1 to 175559)
Worley, K.C.
Direct Submission
Submitted (28-JUN-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
On Jun 26, 2002 this sequence version replaced gi:20335921.
INFORMATION: <http://www.hgsc.bcm.tmc.edu/> or email gc-help@bcm.tmc.edu

COMMENT

CLONE LENGTH: This sequence does not necessarily represent the entire insert of this clone. Overlapping regions of clones are only sequenced and submitted once, so the sequence for the remainder of the insert may be found in the record for the adjacent clones.

Overlapping clones are noted at the beginning and end of the Features listing.

ANNOTATION OF FEATURES:

STSs are identified using ePCR (Genome Res. 7:541-550) searches of a local database that includes entries from dbSTS, GDB, and local mapping efforts.

Repeats are identified using RepeatMasker (A. Smit and P. Green, unpublished.) for Human and Mouse sequences.

Genes and Region of sequence similarity are identified by BLAST (Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-34) to the EST and cDNA sequences. Genes demonstrate at least two exons flanked by consensus splice sites that maintained sequence continuity across the splice junctions. Sequences that are not identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum standard of double strand coverage with a minimum of 2 clones and 2 reads with no ambiguities or 2 chemistries with a minimum of 2 clones and 3 reads with no ambiguities. If the sequence quality for a region does not meet this standard, it will be indicated in the annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality standards - estimated error rate less than 1 per 10,000 bases. Reports of lowest quality individual bases and measures of base quality are listed below. Description of the metrics can be found at URL: <http://www.hgsc.bcm.tmc.edu:8088/quality.info/genbank.annot.attn.html>.

FEATURES

source

Location/Qualifiers

1. 175559
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="X"
/clone="RP11-699A7"

repeat_region

1. 2494
/rpt_family="L1M3f"

misc_feature

1. 2004
/note="overlaps bases 123282..125285 of clone AC121340"

repeat_region

2492..2613
/function="clone overlap"

repeat_region

/rpt_family="L1"

repeat_region

2615..3368
/rpt_family="L1MA8"

repeat_region

3376..3439
/rpt_family="HAL1"

repeat_region

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/rpt_family="HAL1"

repeat_region

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/rpt_family="HAL1"

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repeat_region

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6062..6199
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repeat_region

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6891..6929
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repeat_region

complement(7307..7709)
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repeat_region

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/rpt_family="MIR"

repeat_region

8323..8343
/rpt_family="AT-rich"

repeat_region

8344..8528

COMMENT

Weg 1, D-38124 Braunschweig, Germany, E-mail: info.genome@gbf.de
 On Jan 25, 2002 this sequence version replaced gi:11140934.
 All annotations in this database entry are developed by
 computational tools. It is therefore not explicitly noted in the
 feature lines that evidence is not experimental.
 Mapping was performed at The Sanger Centre
 (cf. <http://www.sanger.ac.uk/HGP/Chr9>)
 Mapping information is available via
<http://webase.sanger.ac.uk/cgi-bin/display?db=acedb9&gprep=179D22>

Center: GBF, Braunschweig
 Center code: GBF
 Web site: <http://genome.gbf.de/>
 Contact: info.genome@gbf.de

----- Project Information
 Center project name:
 Center clone name: bal79D22

----- Summary Statistics
 Sequencing vector: ##;
 Chemistry: Dye-terminator-amersham: 85% of reads
 Chemistry: Dye-primer-amersham: 15% of reads
 Assembly program: Phrap: version 0.990319
 Consensus quality: 0 bases at least Q40
 Consensus quality: 0 bases at least Q30
 Estimated insert size: ##; agarose-fp estimation
 Estimated insert size: 17720; sum-of-contigs estimation

 PROGRAMS AND PARAMETERS USED FOR ANNOTATION:
 + Analysis and annotation were performed with the automatic
 + 'first-pass' annotation and submission tool
 + 'AnnoMitter' (Hornischer & Bloecker).
 + Programs used by 'AnnoMitter':
 + GeneFinder (Green), Vers. 084
 + Organism: human
 + GenScan (Burge & Karlin), Vers. 1.0
 + Used matrix: vertebrate; Minimum score: 0
 + Grail (Xu et al.), Vers. 1.3
 + Organism: human
 + Mzef (Zhang)
 + Prior probability: 0.04; Overlapping number: 0 > Xpound (Thomas
 & Skolnick)

+ Base score cutoff: 0.2; Minimal exon length: 3 bp > 'Repeats':
 BLASTN 2.0.14 (Altschul et al.)
 Database(s): * RepBase: ALU (human), released 22-DEC-1995
 * RepBase: THR ((human), released 22-DEC-1995
 * RepBase: L1 (primate), released 22-DEC-1995
 RepBase: MIR (primate), released 22-DEC-1995
 RepBase: MIR2 (primate), released 22-DEC-1995
 RepBase: THE (primate), released 22-DEC-1995
 Minimum identity: 70 %
 * ESTs: BLASTN 2.0.14 (Altschul et al.)
 Database(s): * emb1 (EST, human), released -DEC-
 (EST), Vers. 67+ (01-JAN-1970) . Using sequence with masked
 repeats
 Minimum score: 60; Minimum identity: 70 %;
 * Tandem Repeats: GDE 2.2 option 'tandem'
 Minimum length 2 bp; Maximum length 20 bp; Score threshold 20
 Treat N's as mismatches? YES; Allow uniform consensi? NO >
 * Inverted Repeats: GDE 2.2 option 'inverted'
 * 'Micro Satellites': GDE 2.2 option 'sputnik' (Abajian) > 'CpG
 Islands': GDE 2.2 option 'cpg'
 CpG island region size 100 bp;
 Minimum GC contents 50 %; Observed/Expected 0.6 > 'STS Scan':
 e-PCR (Schuler)
 Margin: 50; Number of mismatches allowed: 0; Word size: 7
 STS database: 'dbSTS markers'
 * 'tRNA Scan': tRNAscan-SE (Lowe & Eddy), Vers. 1.11.
 Location/Qualifiers

FEATURES

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8789..8973
/rpt_family="MERSA"
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/rpt_family="MLT1B"
10948..11189
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complement(11500..12669)
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13498..13528
/rpt_family="AT-rich"
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/rpt_family="MLT1L"
15438..15671
/rpt_family="MIR"
complement(16593..16897)
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17025..17235
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17868..17980
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17982..20139
/Note="Size of Repeat
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/function="unresolved tandem repeat"
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complement(18163..18193)
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18398..18560
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Query Match      22.8%; Score 41.8; DB 9; Length 175559;
Best Local Similarity 63.4%; Pred. No. 0.0096;
Matches 64; Conservative 0; Mismatches 37; Indels 0; Gaps 0;

52 GAGATCCAAGTCGAGAGAGATGTGGTACGCTATTTATTCCTCCAGTCCTCCCTG 111
||||| | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 75968 GAGATTGAGGACAGAGAGATGATGATATTTGTTCCCTTAGCTACATCCATG 75909

QY 112 CTGGGCTATGGATGACAGCTGGCTGCTACTTCATCTAGAGAAAG 152
| | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 75908 CAGGGCCACTGGTGGCAGTGTCTGCTTTCCTCTAGTAAGG 75868

RESULT 12
HS179D22
LOCUS HS179D22 177220 bp DNA linear PRI 24-JAN-2002
DEFINITION Homo sapiens chromosome 9 BAC RP11-179D22, complete sequence.
ACCESSION AL355433
VERSION AL355433.5 GI:18376585
KEYWORDS HTG.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 177220)
AUTHORS Scharfe,M., Conrad,A., Hornischer,K., Loehnert,T.H., Thies,S. and
Bloeker,H.
TITLE Direct Submission
JOURNAL Submitted (05-MAY-2000) GBF, Dept. of Genome Analysis, Mascheroder

```


Brown, A., Camarata, J., Campopiano, A., Chang, J., Chazaro, B.,
 Choepel, Y., Colangelo, M., Collins, S., Collymore, A., Cook, A.,
 Cooke, P., Dearellano, K., Dewar, K., Diaz, J.S., Dodge, S., Fato, S.,
 Ferreira, P., FitzHugh, W., Gage, D., Galagan, J., Gardyna, S.,
 Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-pierre, N.,
 Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C.,
 Kamat, A., Karatas, A., Kellis, C., LaRocque, K., Lamazares, R.,
 Landers, T., Lehoczy, J., Levine, R., Liu, G., MacLean, C.,
 Macdonald, P., Major, J., Marquis, N., Matthews, C., McCarthy, M.,
 McEwan, P., McKernan, K., Meldrum, J., Meneus, L., Mihova, T.,
 Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C.,
 Norman, C.H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J.,
 Peterson, K., Phunkhang, P., Pierre, N., Pollara, V., Raymond, C.,
 Retta, R., Rieback, M., Riley, R., Rise, C., Rogov, P., Roman, J.,
 Rosetti, M., Roy, A., Santos, R., Schauer, S., Schupback, R., Seaman, S.,
 Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N.,
 Strauss, N., Subramanian, A., Talamas, J., Tesfaye, S., Theodore, J.,
 Topham, K., Travers, M., Travis, N., Triglilio, J., Vassiliev, H.,
 Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.J., Young, G.,
 Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

TITLE JOURNAL

Submitted (24-JAN-2002) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA

On Feb 13, 2002 this sequence version replaced gi:18308592.

All repeats were identified using RepeatMasker:

<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBR

Web site: <http://www-seq.wi.mit.edu>

Contact: sequence_submission@genome.wi.mit.edu

----- Project Information

Center project name: L22007

Center clone name: 640_C_18

 * NOTE: This record contains 73 individual
 * sequencing reads that have not been assembled into
 * contigs. Runs of N are used to separate the reads
 * and the order in which they appear is completely
 * arbitrary. Low-pass sequence sampling is useful for
 * identifying clones that may be gene-rich and allows
 * overlap relationships among clones to be deduced.
 * However, it should not be assumed that this clone
 * will be sequenced to completion. In the event that
 * the record is updated, the accession number will
 * be preserved.

1 679: contig of 679 bp in length
 680 779: gap of 100 bp
 780 1492: contig of 713 bp in length
 1493 1592: gap of 100 bp
 1593 2305: contig of 713 bp in length
 2306 2405: gap of 100 bp
 2406 3042: contig of 637 bp in length
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 3143 3855: contig of 713 bp in length
 3856 3955: gap of 100 bp
 3956 4648: contig of 693 bp in length
 4649 4748: gap of 100 bp
 4749 5465: contig of 717 bp in length
 5466 5565: gap of 100 bp
 5566 6259: contig of 694 bp in length
 6260 6359: gap of 100 bp
 6360 7056: contig of 697 bp in length
 7057 7156: gap of 100 bp
 7157 7810: contig of 654 bp in length
 7811 7910: gap of 100 bp
 7911 8627: contig of 717 bp in length
 8628 8727: gap of 100 bp
 8728 9400: contig of 673 bp in length
 9401 9500: gap of 100 bp
 9501 10214: contig of 714 bp in length
 10215 10314: gap of 100 bp

10315 11020: contig of 706 bp in length
 11021 11120: gap of 100 bp
 11121 11846: contig of 726 bp in length
 11847 11946: gap of 100 bp
 11947 12661: contig of 715 bp in length
 12662 12761: gap of 100 bp
 12762 13467: contig of 706 bp in length
 13468 13567: gap of 100 bp
 13568 14271: contig of 704 bp in length
 14272 14371: gap of 100 bp
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 15070 15169: gap of 100 bp
 15170 15882: contig of 713 bp in length
 15883 15982: gap of 100 bp
 15983 16700: contig of 718 bp in length
 16701 16800: gap of 100 bp
 16801 17501: contig of 701 bp in length
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 19118 19217: gap of 100 bp
 19218 19925: contig of 708 bp in length
 19926 20025: gap of 100 bp
 20026 20753: contig of 730 bp in length
 20756 20855: gap of 100 bp
 20856 21566: contig of 711 bp in length
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 21667 22374: contig of 708 bp in length
 22375 22474: gap of 100 bp
 22475 23181: contig of 707 bp in length
 23182 23281: gap of 100 bp
 23282 24001: contig of 720 bp in length
 24002 24101: gap of 100 bp
 24102 24811: contig of 710 bp in length
 24812 24911: gap of 100 bp
 24912 25600: contig of 689 bp in length
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 26523 27228: contig of 706 bp in length
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 36947 37670: contig of 724 bp in length
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 37771 38479: contig of 709 bp in length
 38480 38579: gap of 100 bp
 38580 39287: contig of 708 bp in length
 39288 39387: gap of 100 bp
 39388 40105: contig of 718 bp in length

* 40106 40205: gap of 100 bp in length
* 40206 40918: contig of 713 bp in length
* 40919 41018: gap of 100 bp in length
* 41019 41723: contig of 705 bp in length
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* 41824 42495: contig of 672 bp in length
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* 43386 44095: contig of 710 bp in length
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* 44895 44994: gap of 100 bp in length
* 44995 45705: contig of 711 bp in length
* 45706 45805: gap of 100 bp in length
* 45806 46517: contig of 712 bp in length
* 46518 46617: gap of 100 bp in length
* 46618 47326: contig of 709 bp in length
* 47327 47426: gap of 100 bp in length
* 47427 48155: contig of 729 bp in length
* 48156 48255: gap of 100 bp in length
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* 49086 49803: contig of 718 bp in length
* 49804 49903: gap of 100 bp in length
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* 53069 53168: gap of 100 bp in length
* 53169 53881: contig of 713 bp in length
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* 53982 54683: contig of 702 bp in length
* 54684 54783: gap of 100 bp in length

Query Match 22.7%; Score 41.6; DB 2; Length 58693;
Best Local Similarity 62.5%; Pred. No. 0.011;
Matches 65; Conservative 0; Mismatches 39; Indels 0; Gaps 0;

OY 59 AAGTGCAGAGGAGGAGTGTGGAGGCTATTATTTATCCCGAGTCCCTCGTGGCT 118
Db 38355 AGGGCAGGAGGAGTGTGGAGGCTTCATTTCTTGTACCTCCCTATTGTGCC 38414
OY 119 ATGATGAACAGTGGCTGACATCTATCAGGAAGAGCTATGCT 162
Db 38415 ATGATTATCAGTCTCTTGTCTCTACCAAGGATAGCTCT 38458

RESULT 14
AP005368/c 111461 bp DNA linear PRI 30-JUL-2002
LOCUS Homo sapiens genomic DNA, chromosome 8q23, clone: KB870F5, complete
DEFINITION sequence.
ACCESSION AP005368
VERSION AP005368.2 GI:22004069
KEYWORDS
SOURCE Homo sapiens pre-pro-B cell cell_line:PLEB 14 - 14 DNA,
clone_lib:Keio BAC library clone:KB870F5.

ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1
Shimizu, N. and Asakawa, S.
AUTHORS
TITLE Homo sapiens DNA chromosome 8 SEQUENCE
JOURNAL Published Only in Database (2002)
REFERENCE 2 (bases 1 to 111461)
AUTHORS Shimizu, N. and Asakawa, S.
TITLE Direct Submission
JOURNAL Submitted (30-MAY-2002) Nobuyoshi Shimizu, Keio University, School
of Medicine, Molecular Biology; 35 Shinnanomachi, Shinjuku-Ku, Tokyo

160-8582, Japan (E-mail: nshimizu@med.keio.ac.jp,
Tel: 81-3-3351-2370, Fax: 81-3-3351-2370)
COMMENT On Jul 30, 2002 this sequence version replaced gi:21280421.
FEATURES
Location/Qualifiers
1. 111461
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="8"
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1. 5368
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TITLE

JOURNAL

REFERENCE

AUTHORS

Submitted (26-DEC-2001) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 3 (bases 1 to 149126)

Birren, B., Linton, L., Nusbaum, C., Lander, E., Ali, A., Allen, N.,
 Anderson, S., Barna, N., Bastien, V., Bloom, T., Boguslavsky, L.,
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REFERENCE

AUTHORS

Submitted (29-APR-2002) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 4 (bases 1 to 149126)

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TITLE

JOURNAL

COMMENT

Submitted (01-MAY-2002) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 On Apr 29, 2002 this sequence version replaced gi:20334653.
 All repeats were identified using RepeatMasker:
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 http://ftp.genome.washington.edu/RM/RepeatMasker.html
 ----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research
 Center code: WIBR
 Web site: <http://www-seq.wi.mit.edu>
 Contact: sequence.submissions@genome.wi.mit.edu
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